



Qy	1	ATGGCAGAAGCTTACCGAATCCAACGGAGATCCCCAATCAAACCTCTTCAG 60	CEY105E8_2	200001	310000
Db	1	ATGGCAGAAGCTTACCGAATCCAACGGAGATCCCCAATCAAACCTCTTCAG 60	CEY105E8_3	300001	410000
Qy	61	CTCTACGTTAAAGCTGCGATTTGCGCATGGCCGATCTTTCTGTCAG 120	CEY105E8_4	400001	510000
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Qy	121	GAATTCTGGATGGAGTTGATGCTCTTATGAGATGGAGTGGAGTGGAGTGAAG 180	CEY105E8_6	600001	702117
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Qy	421	AACGGAAAAAGGAGCCATTGAGGAACTTCAATGTTCAACTCTGAAGGAAACTCAC 480	CEY105E8_16	160001	160000
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Qy	781	CTCCATTGCCCACAAAGGACACAAATCCGAAAGTAGCTTATCGGATATTGTT 840	CEY105E8_28	280001	280000
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RESULT 2		Sequence split into 7 fragments	LOCUS	CEY105E8	Accession AL022594
CEY105E8_2		Fragment Name	Begin	End	
WCOMMENT		CEY105E8_0	1	110000	
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Qy	121	GAATTCTGATGAGTGTGATGCTCTTATGAGATTGGAGTTGGAGTGGAGTCAG	180	
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Qy	241	ATGATTAAGAGGAAALAGAGTGTGATACTACATCTGATATCGAGAGTGTGAG	300	
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Qy	301	TTTCATTTGCAAAAGGAAATTCTGATGTCACATCTTGTAAAGGATCTATCGCT	360	
Db	185255	TTTCATTTGCAAAAGGAAATTCTGATGTCACATCTTGTAAAGGATCTATCGCT	18515	
Qy	361	AGAATAGAGACTGTACAGG	381	
Db	185195	AGAATAGAGAACTGTACAGG	185175	
RESULT	4			
LOCUS	CBRG35124	24174 bp	DNA	linear
DEFINITION	Caenorhabditis briggsae cosmid G35124			INV 04 - NOV - 2000
ACCESSION	AC084558			complete sequence.
VERSION	AC084558.1			
KEYWORDS	HTG.			
SOURCE	Caenorhabditis briggsae			
ORGANISM	Caenorhabditis briggsae			
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida;				
Rhabditidae; Rhabbitidae; Paloderrinae; Caenorhabditis;				
1 (bases 1 to 24174)				
REFERENCE	Washington University Genome Sequencing Center.			
AUTHORS	The C. briggsae Genome Sequencing Project			
TITLE				
JOURNAL				
REFERENCE	2 (bases 1 to 24174)			

AUTHORS	Waterson, R.
TITLE	Direct Submission
JOURNAL	Submitted (04-NOV-2000) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
COMMENT	Submitted by: Genome Sequencing Center Department of Genetics, Washington University, St. Louis, MO 63110, USA e-mail: jspieh@watson.wustl.edu
FEATURES	source
	source
	source
	source
ORIGIN	
NOTICE:	This sequence may not be the entire insert of this clone. It may be shorter because we only sequence overlapping sections once, or longer because we provide a small overlap between neighboring submissions.
LOCATOR/QUALIFIERS	
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Query Match	Score	DB	Length
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Matches 327	80.3%	Pred. No. 1.e-62;	
Matches 327; Conservative	0;	Mismatches 80;	Indels 0;
Qy	0;	Gaps 0;	0;
Db	3448	ATCCGCRGAGAGAGATAAGAGAACTGTACAGGAACCTTCAAACTTCTGCGGAGCAAA	407
Db	23067	ACCCGCAATTAAATTATGATTATTGTTCTGAACTCAAACTCTTCGGGCGCA	23126
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Qy	528	GAGAAAATCTCGATATCTACATTGAAACAGATGACTGAAATACTGTAATGATGCC	587
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Qy	588	ACGTCCTCATCATATTGAAATTGTTGGATTGTGCACTCTTGGATTCCACATAA	647
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Qy	648	TTTCACATCTCTGGCTTATATCCCTCACTGCATACGGATTATTGAGAG	707
Db	23367	TTTCACATCTCTGGAACTCATTCCTACCCGTTACGGTACCGGGGTTCAATTGAGAG	23426
Qy	708	TTGTCGCCGATCGACATATTCACTCATATAAGAACATATGAA	754
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RESULT 5			
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WP2C0MFMENT			
Sequence split into 7 fragments			
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CEY105E8_2		200001	310000
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CEY105E8_5		500001	610000
CEY105E8_6		600001	702117
Continuation (4 of 7) of CEY105E8 from base 300001 (AL022594 Caenorhabditis elega			
Query Match			
Score 238.4%; DB 2; Length 110000;			
Score 27.3%; DB 2; Length 110000;			



JOURNAL	Submitted (08-Oct-2001) Berkeley Drosophila Genome Project, Lawrence Berkeley National Laboratory, One Cyclotron Road, Berkeley, CA 94720, USA	Db	1063 GTTACACATACATCC 1080
COMMENT	Sequence submitted by: Berkeley Drosophila Genome Project		
	Lawrence Berkeley National Laboratory		
	This clone was sequenced as part of a high-throughput process to sequence clones from Drosophila Gene Collection 1 (Rubin et al., Science 2000). The sequence has been subjected to integrity checks for sequence accuracy, presence of a polyA tail and contiguity within 100 kb in the genome. Thus we believe the sequence to reflect accurately this particular cDNA clone. However, there are artifacts associated with the generation of cDNA clones that may have not been detected in our initial analyses such as internal priming, priming from contaminating genomic DNA, retained introns due to reverse transcription of unspliced precursor RNAs, and reverse transcriptase errors that result in single base changes. For further information about this sequence, including its location and relationship to other sequences, please visit our Web site ( <a href="http://fruitfly.berkeley.edu">http://fruitfly.berkeley.edu</a> ) or send email to <a href="mailto:cDNA@fruitfly.berkeley.edu">cDNA@fruitfly.berkeley.edu</a> .		
FEATURES	source		
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	ORIGIN		
	source		
	Query Match 8.8%; Score 76.4; DB 3; Length 1384; Best Local Similarity 52.5%; Pred: No. 4e-09; Matches 167; Conservative 0; Mismatches 151; Indels 0; Gaps 0;		
	Db	495 GCAACTATCCAAATATTGATCAGTTGCTATCCGAGAGAAATTCTCGATATCTACTTGGAAA 554	RESULT 10 AX655393;c
	Db	763 GCAATCGCCGCGATCAAGATCATCTGTCGGCGCAACACCGCCTTCTCACGGCGA 822	LOCUS DEFINITION ACCESSION VERSION
	Db	823 CACCATGCTGCGTGCAGTGTGAGTCTGATCGCAGCATCGTGGCGG 882	Sequence 5263 from Patent WO03000898.
	Db	815 ATGGTCACTTCCTGGATATTGATATTCATCTCGGCTTATATCTC 674	AX655393 AX655393 AX655393.1
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	Db	675 CACTATAAGAACAAATGAAATTGAGAGTATTGCTGGCTATCCGAGCTTCAATGCCAC 734	Oryza sativa Oryza sativa Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Mgnoliophyta; Poales; Poaceae;
	Db	943 TCACTGTAACAGCTGACTTGACGCTTACACATCGTGGCCGCAACATTATCA 1002	Bhrhartoidea; Orzyidae; Orzya.
	Db	735 TCACTATAAGAACAAATGAAATTGAGAGTATTGCTGGCTATCCGAGCTTCAATGCCAC 794	Chang, H.S., Chen, W., Cooper, B., Glazebrook, J., Goff, S.A., Hou, Y.M., Katagiri, F., Quan, S., Tao, Y., Whitham, S., Xie, Z., Zhu, T. and Zou, G.
	Db	1003 TCACTACAAGCTGACAGACTCTCAAATGAAAGAACAGGAGCTGAGACGCCAC 1062	Plant genes involved in defense against pathogens Patent: WO 03000898 A 5/26/2003; Syngenta Participations AG (CH)
	Db	795 AAAAACGACACATTCC 812	FEATURES Location/Qualifiers
	Db	1118 GTTACACATACATCC 1135	
	ORIGIN		
	source		
	Query Match 8.8%; Score 76.4; DB 3; Length 1384; Best Local Similarity 52.5%; Pred: No. 4e-09; Matches 167; Conservative 0; Mismatches 151; Indels 0; Gaps 0;		
	Db	495 GCAACTATCCAAATATTGATCAGTTGCTATCCGAGAGAAATTCTCGATATCTACTTGGAAA 554	RESULT 10 AX655393;c
	Db	763 GCAATCGCCGCGATCAAGATCATCTGTCGGCGCAACACCGCCTTCTCACGGCGA 822	LOCUS DEFINITION ACCESSION VERSION
	Db	823 CACCATGCTGCGTGCAGTGTGAGTCTGATCGCAGCATCGTGGCGG 882	Sequence 5263 from Patent WO03000898.
	Db	815 ATGGTCACTTCCTGGATATTGATATTCATCTCGGCTTATATCTC 674	AX655393 AX655393 AX655393.1
	Db	883 CAAGTACTTGTGCACTTGAATCCGACCGACTTGACGGCCCTGTGGGCTACATGTA 942	GI:29158207 ORGANISM SOURCE ORGANISM SOURCE
	Db	675 CACTATAAGAACAAATGAAATTGAGAGTATTGCTGGCTATCCGAGCTTCAATGCCAC 734	Oryza sativa Oryza sativa Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Mgnoliophyta; Poales; Poaceae;
	Db	943 TCACTGTAACAGCTGACTTGACGCTTACACATCGTGGCCGCAACATTATCA 1002	Bhrhartoidea; Orzyidae; Orzya.
	Db	735 TCACTATAAGAACAAATGAAATTGAGAGTATTGCTGGCTATCCGAGCTTCAATGCCAC 794	Chang, H.S., Chen, W., Cooper, B., Glazebrook, J., Goff, S.A., Hou, Y.M., Katagiri, F., Quan, S., Tao, Y., Whitham, S., Xie, Z., Zhu, T. and Zou, G.
	Db	1003 TCACTACAAGCTGACAGACTCTCAAATGAAAGAACAGGAGCTGAGACGCCAC 1062	Plant genes involved in defense against pathogens Patent: WO 03000898 A 5/26/2003; Syngenta Participations AG (CH)
	Db	795 AAAAACGACACATTCC 812	FEATURES Location/Qualifiers

source	1. .2000 /organism="Oryza sativa" /mol_type="unassigned DNA" /db_xref="taxon:4530"
ORIGIN	<p>Query Match Score 51.2; DB 6; Length 2000; Best Local Similarity 10.4%; Pred. No. 0.018%; Matches 70; Conservative 294; Mismatches 302; Indels 4; Gaps 1;</p> <p>Qy 75 GTCAGGAACTGATGCTGCCGATTCGGCCATTTCTGTCAGGAATTCTGGATGGAA 134 Db 1030 GTGTGAGGTGCTKMRTRSMYTMKRYTKTMTAISSTWKWYAWRSRS 971 Qy 135 GTTGTATGCTCTTATGAGTTGAGTTGAGCTGAGTCAGTGAAGAACGTCACAGTGAA 194 Db 970 RKTWWCTGGKEMATYCGTKMMAAGRWRMWCWYCOMWKWONTSCMWKWKRTWMSWY 911 Qy 195 TTCTGAGGACTTAAGAA---GAACTTCTGGAGCACACCCGATTATGATTGAAG 250 Db 910 TWWGAMRYATYAMRRRWTWKWSWRMWTMKTWAWTWTMCMARWYATGWATWWWW 851 Qy 251 AGGAAAAAGAGCTGACATACACTGATACTGAGAGATTGAGGACGGATCTTCATTTGG 310 Db 850 RYTMYTTCYANTCAKCKYKMANTKWNTTWACWATRWSWRAMAGRWRKXKMRKRAYWR 791 Qy 311 CAAAGGATTCAATGTTCACTCTTGTAAAGGATCCATCGCTGAGAAAGAGAATAGAGA 370 Db 790 WWRCTAGWAKWMSKRYWKWKKVYATRYNNKWMWNSWRWSYRMWSGMGRMRWSA 731 Qy 371 ACTTGTCAGGAACTCAAACTGTTCTGGCAGCAAAAGTAGAGTTCGATAAGGGAAAAA 430 Db 730 WRYCSRMKCAKTYIASARWTKRAKRSYRYRRRWTWKRGTMYTRRYWRCRMTRMSK 671 Qy 431 AGGAGCCATGAGAGTGAAGATCTTCAGCACAGATAAAGGTCACTAAATCGAGTC 490 Db 670 RRKWAGASMSKSCWMMYRGRASWMSKYSKSCAKCCKTRTNTSSYSTMSTGMYMSYYKSMS 611 Qy 491 GTGAGCAACTATCCAAATATGATCAGTTGATATGCTGATATCTGATATCTACTTG 550 Db 610 WTSKNSYMSGMKTCMTYNTSMKGSSTRRSKGMWSGMSRMMKNCRKYMRMKWCWT 551 Qy 551 GAAACAGTATGACTGATAATGACTGTGAACTGATGCCACGCTTCATCATATTGAAATTA 610 Db 550 RRCMCTRWGTMYTTSRSRMNTGTRIKARTTSKRRIMWYKIRKICWYYGMIMCSYMM 491 Qy 611 TTGGATGTCACTTCCTGGATTCTGATTCACATATTGACTCATCTGGCTTATA 670 Db 490 RYGYCRAKCCYAMCWAIAYSGMNNWYKXSKWNRMSKXMSKXGAKG 431 Qy 671 TCCTCACTGATACCGTACAGCAGATTTATGAGGTGTCGCCGATCAGGACATTA 730 Db 430 CYGCKWWTYCYSGYMKWYTNGSYKSRCKYMRMMYKGMIMMMYYSAYSSMMTWYYY 371 Qy 731 TTCACTACTA 740 Db 370 AKYWKWYKR 361</p>
RESULT 11	5 5%; Score 48; DB 9; Length 877;
CR533501	877 bp mRNA linear PRI 22-JUN-2004
LOCUS	Homo sapiens full open reading frame cDNA clone RZP0834B0417D for Gene C1IC4, chloride intracellular channel 4, complete cds, incl. stopcodon.
DEFINITION	
VERSION	GI:49065427
KEYWORDS	Full ORF shuttle clone, Gateway (TM), complete cds.
SOURCE	
ORGANISM	Homo sapiens (human)
ACCESSION	CR533501
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Primates; Cacariniidae; Homo.
REFERENCE	1. bases 1 to 762

Qy	720	TCAGGACATTATCCTACTATAAGACACAATGAAATGATCTGTCACAATCAA	771	
Db	711	TAAGGGCTTGAAATGCTATAGCTATAGTGTAGCCAAAGAGCTCACCAAGTAA	762	
RESULT 12	BC012444	944 bp mRNA linear	PRI 29-JUN-2004	
LOCUS	BC012444	Homo sapiens chloride intracellular channel 4, mRNA (cDNA clone MGC-8812 IMAGE:38861372), complete cds.		
DEFINITION	BC012444			
ACCESSION	BC012444.1	GI:15214635		
KEYWORDS	MGC.			
SOURCE	Homo sapiens (human)			
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
REFERENCE	1 (bases 1 to 944)			
AUTHORS	Straubberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G., Klausner, R.D., Collins, F.S., Wagner, L., Shaenman, C.M., Schuler, G.D., Altschul, S.F., Zeeberg, B.R., Buetow, K.H., Schaeffer, C.F., Bhat, N.K., Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F., Datchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L., Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Tostiyukit, S., Carninci, P., Prange, C., Rana, S.S., Loquellano, N.A., Peters, G.J., Abramson, R.D., Mulayah, S.J., Bosca, S.A., McEwan, P.J., McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S., Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W., Villalon, D.K., Muzny, D.M., Sodergren, B.J., Lu, X., Gibbs, R.A., Faihey, J., Heitton, E., Ketten, M., Madan, A., Madan, A., Rodriguez, S., Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y., Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, B.D., Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M., Butterfield, Y.S., Krzywinski, M.I., Skalska, U., Smalius, D.E., Schenck, A., Schein, J.E., Jones, S.J. and Marrs, M.A.			
TITLE	Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences			
JOURNAL	Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)			
PUBMED	1247932			
REFERENCE	Straubberg, R.			
AUTHORS	2 (bases 1 to 944)			
TITLE	Direct Submission			
JOURNAL	Submitted (15-AUG-2001) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA			
REMARK	NIH-MGC Project URL: <a href="http://mgc.nci.nih.gov">http://mgc.nci.nih.gov</a>			
COMMENT	Contact: MGC help desk Email: <a href="mailto:cgabbs@nih.gov">cgabbs@nih.gov</a>			
	Tissue Procurement: DCTD/DRP			
	cDNA Library Preparation: Life Technologies, Inc.			
	DNA Sequencing by: Baylor College of Medicine Human Genome Sequencing Center			
	Center code: BCM-HGSC			
	Web site: <a href="http://www.hgsc.bcm.edu/cDNA/">http://www.hgsc.bcm.edu/cDNA/</a>			
	Contact: <a href="mailto:ang@bcm.tmc.edu">ang@bcm.tmc.edu</a>			
	Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Lousegod, H., Rowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Nanavati, A.N., Gibbs, R.A.			
FEATURES	source	Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium (LiNL) at: <a href="http://image.llnl.gov">http://image.llnl.gov</a> Series: TRAK Plate: 21 Row: f Column: 8 This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 7330334. Location/Qualifiers 1 .944 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606"		
REFERENCE	1 .944			
AUTHORS				
TITLE				
JOURNAL				
PUBMED				
REFERENCE	1 (bases 1 to 999)			
AUTHORS	Edwards, J.C.			
TITLE	A novel p64-related Cl- channel: subcellular distribution and nephron segment-specific expression			
JOURNAL	Am. J. Physiol. 276 (3 Pt 2), F398-F408 (1999)			
FEATURES	source			
REFERENCE	2 (bases 1 to 999)			
AUTHORS	Edwards, J.C.			
TITLE	Direct Submission			
JOURNAL	Submitted (05-OCT-1998) Medicine, Washington University, 216 S. Kingshighway, St. Louis, MO 63110, USA			
FEATURES	Location/Qualifiers 1 .999			
REFERENCE	1 .999			
AUTHORS				
TITLE				
JOURNAL				
PUBMED				
REFERENCE	1 (bases 1 to 999)			
AUTHORS	Edwards, J.C.			
TITLE	Submitted (05-OCT-1998) Medicine, Washington University, 216 S. Kingshighway, St. Louis, MO 63110, USA			
FEATURES	Location/Qualifiers 1 .999			
REFERENCE	1 .999			
AUTHORS				
TITLE				
JOURNAL				
PUBMED				
REFERENCE	1 .999			
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PUBMED				
REFERENCE	1 .999			
AUTHORS				



COMMENT NEDO human cDNA sequencing project supported by Ministry of Economy, Trade and Industry of Japan; cDNA full insert sequencing: Research Association for Biotechnology (RAB); cDNA library construction: Helix Research Institute (HRI) (supported by Japan Key Technology Center etc.); 5' - & 3'-end one pass sequencing: RAB, HRI, and Biotechnology Center, National Institute of Technology and Evaluation; clone selection for full insert sequencing: HRI and RAB; annotation: HRI and RAB.

FEATURES location/qualifiers

source 1.. .2238

/organism="Homo sapiens"

/mol type="mRNA"

/db\_xref="taxon:9606"

/clone="RHIDPC2003472"

/cell type="dermal papilla cells (HDPC)"

/clone lib="RHIDPC2"

/note="cloning vector: pME18SPL3-primary culture, dermal papilla cells"

ORIGIN

Query Match 5.5\$: Score 48; DB 9; Length 2238;

Best Local Similarity 50.4%; Pred. No. 0.12;

Matches 117; Conservative 0; Mismatches 115; Indels 0; Gaps 0;

Qy	540	ATATCTACTTGGAAACAGTATGACTGATAATGACTGTGAACTGATGCCACGTCTTCATCA	599
Db	637	ATTTCCTGATGGCAATGAAATGACATTAATGCTGATTCGATTCACCTGCTCCAAACTGATAT	696
Qy	600	TATTGGAATTATTGGATTGACTTCTGGATTGATAATTCCACATAATTCACTCATCT	659
Db	697	TGTCAAGGTGGGCCAAAAAAATATGCCAACATTGATATTGATATTCAAAGAAATGACTGGCAT	756
Qy	660	CTGGGTTATACTCTCACTGATACCGTACAGCAGATTATTGAGAGTTTCCGGCGA	719
Db	757	CTGGGATACCTAACATGATACGTTACAGTAGGGACGAGTCACCAATACCTGTCCAGTGA	816
Qy	720	TGAGGACATTATTCATCATAATAAGAACAAATGATCTGTTCAAAATCAA	771
Db	817	TAAGGAGTTGAAATAGCATATAGTGTACGGAAAGACTCACCGATTA	868

Search completed: March 10, 2005, 01:21:16  
Job time : 4126 secs





and DNA pol I. The library was constructed in the lambda Uni-Zap XR vector and has  $1 \times 10^6$  independent recombinants and the average insert size is ~1200 bp. The library was constructed by Sara Lustigman and Michelle Lizotte-Waniowski in the Laboratory of Dr. S. A. Williams. The library is available from Dr. Sara Lustigman (email: [sllustigman@yale.edu](mailto:sllustigman@yale.edu)).





AUTHORS	McCarter, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T., Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y., Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Tsagareishvili, R., Ronko, I., Kennedy, S., McGuire, L., Beck, C., Underwood, K., Stepice, M., Allen, M., Person, B., Swallier, T., Harvey, N., Schurk, R., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R., and Wilson, R.	Db	421 ATTAGATTGGTGGCAAGATTATAGATTAAATCCATAATTACATGTGTA 480
JOURNAL	The Washington Univ. Nematode EST Project, 1999	Qy	661 TGGGGTTATATCCCTACTGCTACGGCATTATTGAGAGTGTCCGGCGAT 720
COMMENT	Unpublished (1999)	Db	481 TGGAAATTATATGCTTACTGCCTATAGCTGCTGATTGCTGATGTGAT 540
CONTACT	McCarter, JP	Qy	721 CAGGACATATTCTACATCTATAAGA 746
THE WASHINGTON UNIVERSITY SCHOOL OF MEDICINE	4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA	Db	541 CAACATTATACACCATACAGAGA 566
RESULT 8			
AI043403	AI043403	AI043403	AI043403
LOCUS	BSBmmfz06110SK	483 bp	mRNA
DEFINITION	Brugia malayi microfilaria cDNA (SAW94LS-BmMf)	linear	EST 01-JUL-1998
ACCESSION	Brugia malayi cDNA clone BSBmmfz06110 5'	mRNA sequence.	
VERSION	AI043403		
KEYWORDS	EST.		
SOURCE	Brugia malayi		
ORGANISM	Brugia malayi		
BRUGIA	Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Pilarioidea; Onchocercidae; Brugia.		
REFERENCE	Williams, S.A.		
AUTHORS	Genes expressed in microfilaria of <i>Brugia malayi</i>		
TITLE	Unpublished (1995)		
JOURNAL	Contact: Steven A. Williams		
COMMENT	Molecular Parasitology Smith College Department of Biological Sciences Department of Biological Sciences, Clark Science Center, Smith College, Northampton, MA, 01063, USA Tel: 4135853826 Fax: 4135853786		
EMAIL	genome@smith.edu		
EST	Submitted by Molecular Parasitology Group, New England Biolabs, Inc., 32 Tozer Road, Beverly, MA, 01915, USA. Email: dnaseq@neb.com		
dnaseq@neb.com	Site in the clone designation refers to 'Subtracted Method Z'. The numerical designation after the SZ refers to the microtiter tray number (01 - 48) followed by the letter and number of the microtiter tray position. Method 'Z' refers to sequenced clones which have been selected after hybridization subtraction (18,000 mass-excised colonies gridded as a high density array on nylon filters). Colonies not represented in the grid were used as templates for the sequencing reactions.		
SEQ PRIMER	Seq primer: pBluescript SK.		
LOCATION/QUALIFIERS	Location/Qualifiers		
1.	1. .367		
LOCATION/QUALIFIERS	/organism="Strongyloides stercoralis"		
1.	/strain="Filariiform larvae obtained from humans"		
1.	/mol_type="mRNA"		
1.	/db_xref="Taxon:6248" (Stratagene)"		
1.	/lab_host="XL-1 Blue MRF" (Stratagene)"		
1.	/clone_id="TB95TM-SSEH"		
1.	/clone="Vector: Lambda Uni-ZAP XR (Stratagene): Site 1: EcoRI; Site 2: XbaI; mRNA was purified from 4 x 10 <sup>35</sup> Filariiform larvae which had been isolated from infected humans. cDNA was constructed and, using adaptors, was cloned unidirectionally into the vector from the EcoRI site to the XbaI site. The library has an unamplified titer of 1.5 x 10 <sup>86</sup> pfu/ml and an amplified, undiluted titer of 7 x 10 <sup>89</sup> pfu/ml. The average insert size of the unamplified library is 975 bp (range, 500-1500)."		
ORIGIN			
FEATURES			
source			
1.	1. .483		
1.	/organism="Brugia malayi"		
1.	/mol_type="mRNA"		
1.	/strain="TR5 Labs"		
1.	/db_xref="Taxon:6279"		
1.	/clone="BSBmmfz06110"		
1.	/lab_host="XL-Blue MRF"		
1.	/clone="Brugia malayi microfilaria cDNA (SAW94LS-BmMf)"		
1.	/note="Vector: lymphatic filarial parasite of humans."		
1.	/note="lambda Unizap XR; Site 1; Ecor I; Site 2: Xba I; isolated from microfilariae of Brugia malayi from jirds and converted to double stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNAPol I. The library had 3.5 x 10 <sup>85</sup> independent recombinants and average insert size was 900 base pairs. The library was constructed by Lori Saunders. The library is available from Dr. S.A. Williams, email genome@smith.edu."		
ORIGIN			
Query Match			
Best Local Similarity			
Matches			
348	23.3%; Score 203.6; DB 2; Length 567;		
0	Pred. No. 3.9e-48;		
0	Mismatches 209; Indels 9; Gaps 1;		
Qy	190 GTGAAATTCTGAGCATTAAGAGAACTTCTGGAGCACACCCGATTATGATTGAA 249		
Db	1 GTTAATTCTGAGCTTAAATTTCTGGAGCTACCTCTTGTGAAAGATTATTGATTGAA 60		
Qy	250 GAGGAAGAGACCTGACATACACTGATACTGAGATGAGGAGGATCTTCATTC 309		
Db	61 GTTCCAAAGATGCAACATATTGAGATAAGATGAAATGAGATGAAATGATCTT 120		
Qy	310 GCAAGAAATTCAATGTTCCACTCTTGTAAAGGATCCATCGCTGAGAGAGATAAG 369		
Db	121 GCCAAAGAAATTGTGATGCTCTTGTAAAGATATTACTGTGAAAGATTATTGCA 180		
Qy	370 AACTGTACAGAACACTCAACTGTTCTGGCAGAAAGTGTGAGTGTGAAAGAA 429		
Db	181 TCACTTATTAGAAATTAAATTATTCTTAAGCAAAACTGATCATATAAGTAAAG 240		
Qy	430 AAG-----GAGCCATCGAGGTGAAATCTCCAGCACATTAAGTTCACTTCAC 480		
Db	241 AGATAAAATAGAAATAATACTTTACAGAAAGTTTACACATCTGTGTTGTCCT 300		
Qy	481 AATCGACTCTGAGAACATCTGAAATATGATCAGTGTGCTATCGAGAAAAATCTCGA 540		
Db	301 ACAAAATTACTAGAAATAGCATATTGATCAATGATTTAGCAAGAGGTCAAGA 360		
Qy	541 TATCTACTGTGAAAGCTGAACTGAAATGACTGTGAACTGCTGCAAGTCTCATCT 600		
Db	361 TATTATTATCACAAATTGATGTTGAGTGTGATGATGATGATGATCATCT 420		
Qy	601 ATTGAATATTGGATTGTCACITCTCTGGATTGATTCACATCTCACATCTC 660		
Query Match	23.1%	Score 202;	DB 1;
Best Local Similarity	75.9%	Length 483;	
Matches	274;	Conservative 0;	
Qy	361 TATTATTATCACAAATTGATGTTGAGTGTGATGATGATGATCATCT 420	Indels 4;	Gaps 2;

Qy	45	AAAACCTCTCTGAGCTCTAGTAAGCGATTGATGGCTGCCCATGGGC	104	FEATURES	The vector to vector length is 632
Db	125	AAAACCGGTTCTGAGCTTCTGAGCTTCTGAGAAATCTGGATCGAA	184	source	Seq primer: Sp6.
Qy	105	CGATCTTPTCTGAGATTCTGAGTGTGATGCTTATGAACTGGATTG	164	Location/Qualifiers	Location/Qualifiers
Db	185	ATGTTGTTGTTGCAAGATTCTGGATGAACTTACGCTCTCATGAA	244		1. 566 /organism="Meloiodyne paraensis" /mol_type="mRNA" /db_xref="taxon:189293" /tissue_type="whole organism" /dev_stage="egg" /lab_host="DH10B" /clone_id="Meloiodyne paraensis egg SMART pgEM" /note="Vector: pGEM-11zf(+)" (Promega); Site 1: XhoI; Site 2: NotI; Cloned unidirectionally. Poly(A)+ RNA was concentrated and purified using Dynabeads (Dynal) and mRNA was eluted for first strand synthesis. First strand cDNA was created using MMLV RT (PowerScript, Clontech) and primed with oligo(dT) with XhoI site (primer CDSIII/3-XhoI) and 5' SMART anchor, added using chimeric DNA-RNA oligo (SMART-NotI-r-GGG). 12 PCR cycles were done using first strand and primers specific to SMART oligo (5' PCR primer) and 3', end (XhoI-No-dT). Double stranded cDNA was digested using XhoI/NotI, fractioned on Chroma-spin 400 columns (Clontech) and ligated to digested pGEM-11zf(+) plasmid. Chemically competent DH10B cells were used as host cells. Library constructed by Joanna Carlson."
Qy	165	ACGAGTCGAACTGAACTGCACTGAACTTCTGAGCATTAAAGGA	223		
Db	245	TGAGTGAAGTGAACAGTAAACATCTGAGCATTAAGAGTTCCTGG	304		
Qy	224	GAGCACACCACGGATTATGTTGAGGAAAAAGAGCTGACATAC	283		
Db	305	GACCAACACCACGGATTATGGTAAACAAAGATG -- -CACTTATA	361		
Qy	284	AGATTGAGGAGGAGCTTCTATGGAAAGGAATTCAAGTGTGAAAGG	343		
Db	362	AAATTGAGGAGGCAATTTCATTTGAGAAAAGNTCAATGTGCCA	421		
Qy	344	ATCCATCGCTGCTGAGAACTTCAACTGTCTCCGAG 403			
Db	422	ATCCAGTGTGNGAAAGATGAACTTCAAACTTTAGT 481			
Qy	404	C 404		ORIGIN	
Db	482	C 482			Query Match 21.6%; Score 189; DB 7; Length 566;
					Best Local Similarity 67.3%; Pred. No. 7.4e-44;
					Matches 267; Conservative 0; Mismatches 130; Indels 0; Gaps 0;
Qy					Qy 440 CGAGAGTTGAGATCTTCTCCAGCAGTAAAGTCACACTAACATCGAGTCAGTCTGTGAGCAAC 499
					Db 38 CAATAGAGAGGGCCCTCCCCACAGTGACATCTCACAGATAAATTATTGGAGCAAT 97
Qy					Qy 500 TATCCAAATATGATGAGTGTGTGTATCGAGAAAAAATCTCGATATATCTACTTGAAACAGTA 559
					Db 98 TGGCCAAATATGACAGTGTGGCTGATCGTTCTCTCGTTATCTAAGTGTAGTCAGTCA 157
Qy					Qy 560 TGACTGAAATGACTGTGAACTGACATCATATGGATTATTGGATTGTG 619
					Db 158 TGACTGAAATGACTGTGAGATTGAGATTGACCATTTGACCAATTCGATATTGGAGAAC 217
Qy					Qy 620 CACTCTGTGATTGATTCACATATTTCACATCTCTGGCTTATATCCTCACTG 679
					Db 218 GCCTTTGAATTTTACATTCCACCTTCACTTATGGCATATATATACTG 277
Qy					Qy 680 CATACTGTACAGCAGATTATTGAGATGTTGGCTCCCGATGGGACATTATTCACT 739
					Db 278 CATATGGAACAGCTGGTTTATTGAACTCTGTCGGCTGTGATCAAGATATTGCACTT 337
Qy					Qy 740 ATAAGGACAAATGAACTGTGCAAACTAACATCGCTAACAAAA 799
					Db 338 ATAAGGACAAATTAACCTGGCAATTGATGGAATGACTTGAGCACCAAGAA 397
Qy					Qy 800 CGCACACAAATCCGGAAAAAGTGTATGGATATTGTC 836
					Db 398 CATTACCAATTCCAGAAGATTTCTCAAGATATACG 434
				RESULT 10	
				CB374401	CB374401
				LOCUS	LOCUS
				DEFINITION	DEFINITION
				CB374401	using rup9h02_Y1 Heterodera glycines
				ACCESSION	ACCESSION
				CB374401	CDNA 5', similar to WP:CE25703 Y105E8C.A.;
				VERSION	mRNA sequence.
				CB374401.1	GI:29049758
				KEYWORDS	EST.
				ORGANISM	Heterodera glycines
				COMMENT	Eukarya; Metazoa; Nematoda; Chromadorea; Tylenchida; Heterodera; Glycines
				COMMENT	Tylenchoidea; Heteroderidae; Heterodrinae; Heterodera; Tylenchina;
				COMMENT	The Washington Univ. Nematode EST Project, 1999
				COMMENT	Washington University School of Medicine
				COMMENT	4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
				COMMENT	Tel: 314 286 1800
				COMMENT	Fax: 314 286 1810
				COMMENT	Email: est@watson.wustl.edu
				COMMENT	Cloned unidirectionally. Poly(A)+ RNA was concentrated and purified
				COMMENT	using Dynabeads (Dynal) and mRNA eluted for first strand synthesis.
				COMMENT	First strand cDNA was created using MMLV RT (PowerScript, Clontech)
				COMMENT	and primed with oligo(dT) with XhoI site (primer CDSIII/3-XhoI) and
				COMMENT	5' SMART 'anchor' added using chimeric DNA-RNA oligo
				COMMENT	(SMART-NotI-r-GGG). 12 PCR cycles were done using first strand and
				COMMENT	primers specific to SMART oligo (5' PCR primer) and 3'
				COMMENT	end (XhoI-No-dT). Double stranded cDNA was digested using XhoI/NotI,
				COMMENT	Fractionated on Chroma-spin 400 columns (Clontech) and ligated to
				COMMENT	digested pGEM-11zf(+) plasmid. Chemically competent DH10B cells
				COMMENT	were used as host cells. Library constructed by Joanna Carlson of
				COMMENT	North Carolina State University.
				COMMENT	Putative full length read

AUTHORS	McCarter, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T., Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Gibbons, M., Ritter, P., Bennett, J., Franklin, C., Tsagareishvili, R., Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe, M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R., and Wilson, R.	Qy 404 CAAACTAGACTTCCGATAAGGAAAAAAGGAGCCATCGAGAGTCAGAGATCTTCAGCAC 463 Db 406 CCAAAACGAATTGGCAARGCAGCAGAACCCAGTCATTGAAAGCTTCGGCCCC 465
TITLE	The Washington Univ. Nematode EST Project, 1999	Qy 464 A 464 Db 466 A 466
JOURNAL	Unpublished (1999)	
COMMENT	Contact: McCarter, JP The Washington Univ. Nematode EST Project, 1999 Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800 Fax: 314 286 1810 Email: est@watson.wustl.edu	RESULT 1.1 BT741942 LOCUS 396 bp mRNA linear EST 25-SEP-2001 DEFINITION kt3b07_Y1 Strongyloides ratti L1 pAMP1 v3 Chiapelli McCarter Strongyloides ratti cDNA 5, similar to WP:CE2511 Y10568R_Y ;contains element MER1.9 repetitive element ;, mRNA sequence. BT741942.1 GI:15742898
FEATURES	source /clone lib="Heterodera glycines" /note="Vector: pBlueScript SK- (Stratagene), Site 1: XbaI; Site 2: EcoRI; This library was generated by cloning cDNAs directionally into Uni-ZAP(Stratagene) (M3 primer/EcoRI are at the 5'-end and T'XbaI are at the 3'-end). The library was excised (now in pBlueScript SK+) and normalized (Bonaldo et al 1996 Genome Research 6:791-806). Library constructed by Thomas Baum (tbbaum@iastate.edu), Iowa State University, Plant Pathology Department and Jeff McDermott (jpmcderm@iastate.edu)."	ACCESSION BT741942 VERSION EST KEYWORDS Strongyloides ratti Strongyloides ratti Panagrolaimoidea; Strongyloidae; Strongyloides. ORGANISM Bacteria; Metazoa; Nematoda; Chromadorea; Rhabditida; Panagrolaimoidea; Strongyloidae; Strongyloides. AUTHORS McCarter, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Theising, B., Wylie, T., Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y., Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Tsagareishvili, R., Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe, M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R., and Wilson, R.
FEATURES	source /clone lib="Heterodera glycines virgin female" /note="Vector: pBlueScript SK- (Stratagene), Site 1: XbaI; Site 2: EcoRI; This library was generated by cloning cDNAs directionally into Uni-ZAP(Stratagene) (M3 primer/EcoRI are at the 5'-end and T'XbaI are at the 3'-end). The library was excised (now in pBlueScript SK+) and normalized (Bonaldo et al 1996 Genome Research 6:791-806). Library constructed by Thomas Baum (tbbaum@iastate.edu), Iowa State University, Plant Pathology Department and Jeff McDermott (jpmcderm@iastate.edu)."	REFERENCE 1 (bases 1 to 396) TITLE Unpublished (1999) JOURNAL COMMENT Contact: McCarter, JP The Washington Univ. Nematode EST Project, 1999 Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800 Fax: 314 286 1810 Email: est@watson.wustl.edu
FEATURES	source /clone lib="Heterodera glycines virgin female" /note="Vector: pBlueScript SK- (Stratagene), Site 1: XbaI; Site 2: EcoRI; This library was generated by cloning cDNAs directionally into Uni-ZAP(Stratagene) (M3 primer/EcoRI are at the 5'-end and T'XbaI are at the 3'-end). The library was excised (now in pBlueScript SK+) and normalized (Bonaldo et al 1996 Genome Research 6:791-806). Library constructed by Thomas Baum (tbbaum@iastate.edu), Iowa State University, Plant Pathology Department and Jeff McDermott (jpmcderm@iastate.edu)."	FEATURES Source 1. 396 "Strongyloides ratti" /organism="Strongyloides ratti" /mol_type="mRNA" /db_xref="Taxon:34506" /dev_stage="L1" /lab_host="DH10B" /clone_lib="Strongyloides ratti L1 pAMP1 v3 Chiapelli McCarter" /note="Vector: pAMP1 (Gibco); The library was constructed by Brandi Chiapelli and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dynal). PCR based library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UBC sites of pAMP1. Nematodes were provided by Dr. Mark Viney of Bristol, UK."
ORIGIN		Query Match 21.3%; Score 185.8; DB 6; Length 480; Best Local Similarity 65.1%;保守性 0; Mismatches 147; Indels 0; Gaps 0;
FEATURES		Query Match 21.3%;保守性 0; Mismatches 147; Indels 0; Gaps 0;
Query	44 CAAACCTCTCTCGAGCTCTAGTAAAGGTCAGGAATGATGCTCGCGCATGGAG 103	FEATURES Source 1. 396 "Strongyloides ratti" /organism="Strongyloides ratti" /mol_type="mRNA" /db_xref="Taxon:34506" /dev_stage="L1" /lab_host="DH10B" /clone_lib="Strongyloides ratti L1 pAMP1 v3 Chiapelli McCarter" /note="Vector: pAMP1 (Gibco); The library was constructed by Brandi Chiapelli and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dynal). PCR based library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UBC sites of pAMP1. Nematodes were provided by Dr. Mark Viney of Bristol, UK."
Query	46 CAAAGCCCTTGTCTGAGTTCAGGAACTCTGATGGGGTTGATGCTCTTATGAGATTGGATGTG 105	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	104 CGGATCTTTCTGTCAGGAATCTGGATGGGGTTGATGCTCTTATGAGATTGGATGTG 163	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	106 CATGCTTATTTGCCCAGAAATTGGAACTCTAGCTTCTAGAAGTGGCTGTC 165	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	164 CACGAGTGAAGTGAACAATGTCGAACTTCAAGGAAATTCTGTCAGCTTTCGG 223	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	166 TTCTGTGTGAGTAAACCTGTAATTCGTCATTCGAAAGTCAAAAGACTTTCTGG 225	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	224 GAGCCACACCACCGATAATGATTGAAGGAAAAAGCCTGACATACACNGATAATCGAG 283	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	226 GAGCACGCCGCAATATGGGAACTGGCAAGAACGAACTGGACAACTCG 285	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	284 AGATGGAGGAGGATTTCTTCAATTGGCAAGGAAATTCTGTCAGCTTTCGG 343	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	286 ACATCGAGCAGGCAATTCCACATTTGCAAGAACTCTGTGCCATTCTTCGAGAC 345	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	344 ATCCATCGCTTGAAGGAGTAAAGCTGAACTCAATGAGCTTCTGAGCAACTATCCAA 403	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Query	346 ATCCGGAAATGGCCAAAGAACCATCCAGGATTGAGAACTTCCTCCAAAG 405	Query Match 20.2%; Score 176.4; DB 4; Best Local Similarity 66.1%;保守性 0; Mismatches 131; Indels 0; Gaps 0;
Db	7 GAACCTCPACCAATCTGCTGAGCTGTGTCGATCAAAATTGTTAGAACATTGGCATCA 66	Db 7 GAACCTCPACCAATCTGCTGAGCTGTGTCGATCAAAATTGTTAGAACATTGGCATCA 66

Query	Subject	Sequence
Qy	508 ATTGATCAGTGTCTGATCCGAGAAAATCTCGATATCTGAAACGTTGACTGAA 567	118 CATGCTTATTGCAAGAAGATTTCGATGCAACTCTACGCTTCTAGAAGTGGCTG 177
Db	67 ATTGACAATTGTTAGCAAAAGGTTAAGAATTGTTATCAAAAGTTATGTAA 126	164 CACGAGTCGAGTGAAGACTGTCACTGAACTGGAATTCTGAAACACTTCTCG 223
Qy	568 TATGACTGAACTGATGGCAGCCTTCATCATATTGAAATTGATTGATCTCT 627	178 TTGCTGTTGAGTTAAACTGTGATGTCGATTCGAACTTGGCTTTCG 237
Db	127 TATGTTGAGTTAATGCAAGATTACATCATATTGAAATTGTTGGCAAAAGTTATA 186	Qy 224 GAGCACAACCCGATTTGATGAGAAGGAGCTGACATACGTATAATCGAG 283
Qy	628 GATTGATTCAGATATTCACTCATCTCGGCTTATCCTCACTGATACCGT 687	Db 238 GAGCAGCGCCATAATGGAAAGAAGAGCGCAATGGCAAACTGAGCAG 297
Db	187 GATTGTAATTCAGATATTCACTCATCTCGGCTTATCCTCACTGATACCGT 246	Qy 284 AGATGAAAGCAGGATCTTCAATTGCAAGGAAATCATATGTCACCTTTGAAAGG 343
Qy	688 ACAGCAGGATTTATTGAGGTTGTCGGCATGGACATTATTCACTATAAAGA 747	Db 298 ACATGAGCCGACGATTTCACACTTGCACCAATTCCATGTCGCACTTTCGAAAGG 357
Db	247 ACAGTGCTTTATTGAAAGTTGCGCAGTGATGATGAGATACCTTAAAGGA 306	Qy 344 ATCCATCGGTGAGAAGAGATAAGAACTGTGACAGAACCTTAAACGTGCGAG 403
Qy	748 CAAATGAAATTGTCACAAATCACACGTTGAAACCCCTCCATGGCAACAAAGCACCA 807	Db 358 ATCCGGAAATGGCAAGGACATCCGAGGACTTGTGAGGAACTTGTGAG 417
Db	307 CAATTAATATTTCATCACTAACGTGAACATTACGACCAACAAATAACACA 366	Qy 404 CAAAGTAGAGTTGC 418
Qy	808 ATTCCGGAAAAGGCTATGGATAT 833	Db 418 CAAAAGGAAATTG 432
Db	367 ATTCCAAGGGCTTATGAAAT 392	
		RESULT 1.3
		CA868923
	LOCUS	CA868923
	DEFINITION	567 bp mRNA linear EST 20-DEC-2002
	ACCESSION	pw91906.y1
	VERSION	Haemonchus contortus
	KEYWORDS	EST
	ORGANISM	Haemonchus contortus
	REFERENCE	Eukaryota; Nematoda; Chromadorea; Rhabditida; Strongylida; Trichotrichyloidea; Haemonchidae; Haemonchines; Haemonchus. (bases 1 to 567)
	AUTHORS	McCarrier, J., Clifton, S., Chiappelli, B., Pape, D., Martin, J., Wylie, T., Dante, M., Marra, M., Hillier, L., Kucaba, T., Thausing, B., Bowers, Y., Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Tsagareli, Shvili, R., Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe, M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and Wilson, R.
	TITLE	Unpublished (1999)
	JOURNAL	Contact: McCarter, JP
	COMMENT	The Washington Univ. Nematode EST Project, 1999
		Washington University School of Medicine
		4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
		Tel: 314 286 1800
		Fax: 314 286 1810
		Email: est@watson.wustl.edu
	FEATURES	The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dynal). PCR based library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UDG sites of PAMP1. Intestinal RNA was provided by Dr. Douglas Jasmer of Washington State University (djasmer@vetmed.wsu.edu).
	source	Intestinal RNA was provided by Dr. Douglas Jasmer of Washington State University (djasmer@vetmed.wsu.edu).
		Seq primer: -40R from Gibco
		High quality sequence stop: 410.
		Location/Qualifiers
		1. .450
		/organism="Haemonchus contortus"
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		/db_xref="taxon:6289"
		/lab_host="DH10B"
		/clone_lib="pAMP1; Site 1: NotI; Site 2: SalI; The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dynal). PCR based
Qy	44 CAAACCTTCTGAGCTTACCTAAAGCTGATGAAATTGATGTCCTCCGATTTGGAG 103	
Db	58 CAAAGCCCTTGTGCTGAGTTATGAAAGCCCTCAGGATGACGCCAGCATCGGTG 117	
Qy	104 CGATCTTCTGAGGATTCTGATGAGTGGATGATGCTTATGAGATTGAGTGG 163	

library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UDG sites of pANP1. Intestinal RNA was provided by Dr. Douglas Jaeger of Washington State University (djaeger@vetmed.wsu.edu)."

## ORIGIN

Query Match 17.5%; Score 152.6; DB 6; Length 567;  
Best Local Similarity 69.8%; Pred. No. 3, 4e-32; Mismatches 0;  
Matches 206; Conservative 0; Indels 0; Gaps 0;

Qy 559 ATGACTGAAATGACTGTGAACGTGACGTCTTCATATTGAGATG 618  
Db 4 ATGACGGAAATGATGTTGAACCTTATGCCAGACTAGTCATCGGCCAG 63

Qy 619 TCACTCTTGGATTCGATATTGATTCATCATCTGGGTATATCCCTCACT 678

Db 64 AGAATGCTCAATTGATTTCGATTTCCTCGCAATTAACTTAACTGACC 123

Qy 679 GATACCCGTACAGGAGTATTGAGATGTCGCCCGATCAGGACATTATCAC 738

Db 124 GCCTATCAGGCGCGCCCTCATGGAGCTGTCGCCGATCAAGATACTGATCAT 183

Qy 739 TATAAGGACAATGAAATGAACTGTGTCAAAATACGTGAAACCTCCAAATGCCAAGAAA 798

Db 184 TATAAGGACAATTAATGTTACAAATACGTGAACTGCTGAGCTCGCAAGAG 243

Qy 799 ACCGACAGAAATTCCGGAAAATGGCTATGGATATTCGTTAAAGACTTGTC 853

Db 244 ACCATACCACTCCGGAAAGGGTGTGATGGACATGAAAGAACTGGCTGATC 298

RESULT 14

BS225023 BG225023 426 bp mRNA linear EST 09-MAY-2001

DEFINITION to SW:CL:2 HUMAN O15247 CHLORIDE INTRACELLULAR CHANNEL PROTEIN 2, ,

MRNA sequence.

BG225023

EST.

KEYWORDS Strongyloides stercoralis

SOURCE Strongyloides stercoralis

ORGANISM Strongyloides stercoralis

REFERENCE 1 (bases 1 to 426)

AUTHORS McCarter,J., Clifton,S., Chiappelli,B., Pape,D., Martin,J.,

Wylie,T., Dantes,M., Marra,M., Hillier,L., Kucaba,T., Theising,B.,

Bowers,Y., Gibbons,M., Ritter,E., Bennett,J., Franklin,C.,

Tsagareishvili,R., Ronko,I., Kennedy,S., Maguire,L., Beck,C.,

Underwood,K., Stepco,M., Allen,M., Person,B., Swaller,T.,

Harvey,N., Schurk,R., Kohn,S., Shin,T., Jackson,Y., Cardenas,M.,

McCann,R., Waterston,R., and Wilson,R.

TITLE Unpublished (1999)

COMMENT Contact: McCarter, JP

The Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA

Email: est@watson.wustl.edu

The library was constructed by Dr. Thomas Nutman and colleagues of

NIAD, NIH (trutman@nih.gov). DNA Sequencing by: Washington

University Genome Sequencing Center St. Louis.

Seq primer: -40P from Gibco

High quality sequence stop: 418.

Location/Qualifiers 1..426

/organism="Strongyloides stercoralis"

/mol type="mRNA"

/strain="Filariform larvae obtained from humans"

/db\_xref="taxon:6248"

/lab host="XL-1 Blue MRF" (Stratagene)"

/clone lib="TBN95TM\_SSST"

/note="Vector: Lambda Uni-ZAP XR (Stratagene); Site 1: EcoRI; Site 2: XbaI; mRNA was purified from 4 x 10<sup>8</sup> filariform larvae which had been isolated from infected humans. cDNA was constructed and, was cloned unidirectionally into the vector from the EcoRI site to the XbaI site. The library has an unamplified titer of 1.5 x 10<sup>6</sup> pfu/ml and an amplified, undiluted titer of 7 x 10<sup>9</sup> pfu/ml. The average insert size of the unamplified library is 975 bp (range, 500-1500)."

ORIGIN

Query Match 146.8%; Score 146.8; DB 4; Length 426;

Best Local Similarity 70.5%; Pred. No. 1.6-31; Mismatches 0; Indels 0; Gaps 0;

Matches 196; Conservative 0; Mi matches 82;

Query Match 16.8%; Score 146.8; DB 4; Length 426;

Best Local Similarity 70.5%; Pred. No. 1.6-31; Mismatches 0; Indels 0; Gaps 0;

Matches 196; Conservative 0; Mi matches 82;

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Best Local Similarity 70.5%; Pred. No. 1.6-31; Mismatches 0; Indels 0; Gaps 0;

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Matches 196; Conservative 0; Mi matches 82;

Query Match 16.8%; Score 146.8; DB 4; Length 426;



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Grange Bléneau (uspto)

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ON nucleic - nucleic search, using SW model									
Run on:	March 9, 2005, 23:54:49 ;	Search time 194 Seconds							
(without alignments)									
7363.245 Million cell updates/sec									
Title:	US-10-612-379-1								
Perfect score:	873	1 atggcagaatcttaccaat.....ccgatgttaatgttcattaa	873						
Sequence:									
Scoring table:	IDENTITY_NUC								
Gapcost:	Gapext 1.0								
Searched:	1202784 seqs, 81813359 residues								
Total number of hits satisfying chosen parameters:	2405568								
Minimum DB seq length:	0								
Maximum DB seq length:	2000000000								
Post-processing:	Minimum Match 0%								
	Maximum Match 100%								
	Listing first 45 summaries								
Database :	Issued Patents NA:*								
	1: /cgn2_6/ptodata/1/ina/5A_COMB.seq:*								
	2: /cgn2_6/ptodata/1/ina/5B_COMB.seq:*								
	3: /cgn2_6/ptodata/1/ina/5A_COMB.seq:*								
	4: /cgn2_6/ptodata/1/ina/6B_COMB.seq:*								
	5: /cgn2_6/ptodata/1/ina/PCTUS_COMB.seq:*								
	6: /cgn2_6/ptodata/1/ina/backfiles..seq:*								
Pred. No.	is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.								
SUMMARIES									
Result No.	Score	Query Match	Length	DB ID	Description				
1	76.4	8.8	1090	4	US-09-210-767-26167	Sequence 26167, A			
2	76.4	8.8	1090	4	US-09-210-767-10712	Sequence 10712, A			
c	3	47.4	5.4	7218	1	US-08-232-463-14	Sequence 14, Appl		
c	4	46.4	5.3	819	3	US-08-792-014-2	Sequence 2, Appl		
c	5	46.4	5.3	819	3	US-09-443-948-2	Sequence 2, Appl		
c	6	46.4	5.3	819	4	US-09-630-196-2	Sequence 2, Appl		
c	7	43.4	5.0	1652	4	US-09-630-312D-815	Sequence 815, Appl		
c	8	43.4	5.0	1689	4	US-09-643-051-780	Sequence 5780, Appl		
c	9	4.1	4.7	1068	4	US-09-710-794-3	Sequence 3, Appl		
c	10	39.8	4.6	601	4	US-09-949-016-107953	Sequence 107953,		
c	11	39.8	4.6	601	4	US-09-949-016-107954	Sequence 107954,		
c	12	39.8	4.6	39090	4	US-09-949-016-14720	Sequence 14720, A		
c	13	39.2	4.5	1529	4	US-09-533-029-49	Sequence 49, Appl		
c	14	38.4	4.4	1141	4	US-09-612-408-22	Sequence 22, Appl		
c	15	38.4	4.4	253345	4	US-09-949-016-12656	Sequence 12656, A		
c	16	38.4	4.4	253364	4	US-09-949-016-13639	Sequence 13639, A		
c	17	37.8	4.3	832	4	US-09-621-767-2813	Sequence 2813, Appl		
c	18	37	4.2	14066	4	US-09-601-198-56	Sequence 56, Appl		
c	19	36.8	4.2	1664976	4	US-08-916-421B-1	Sequence 1, Appl		
c	20	36.8	4.2	1664976	4	US-09-612-570-1	Sequence 1, Appl		
c	21	36	4.1	42381	4	US-09-949-016-12012	Sequence 12012, A		
c	22	36	4.1	168394	4	US-09-949-016-13002	Sequence 13002, A		
c	23	36	4.1	183770	4	US-09-949-016-15494	Sequence 15494, A		
c	24	35.8	4.1	95561	4	US-09-949-016-12768	Sequence 12768, A		
c	25	35.8	4.1	95561	4	US-09-949-016-13306	Sequence 13306, A		
c	26	35.6	4.1	95561	4	US-09-949-016-1307	Sequence 161307, A		
c	27	35.6	4.1	601	4	US-09-949-016-86515	Sequence 86515, A		
c	28	35.6	4.1	183770	4	US-09-949-016-14597	Sequence 14597, A		
c	29	35.6	4.1	601	4	US-09-949-016-14596	Sequence 14596, A		
c	30	35.4	4.1	832	4	US-09-621-976-2813	Sequence 15753, A		
c	31	35.4	4.1	236964	4	US-09-949-016-15753	Sequence 13294, A		
c	32	35.4	4.1	250715	4	US-09-949-016-13294	Sequence 141597, A		
c	33	35.4	4.0	601	4	US-09-949-016-14597	Sequence 17, Appl		
c	34	35	4.0	1497	3	US-09-232-468A-17	Sequence 14, Appl		
c	35	35	4.0	1497	3	US-09-784-984B-14	Sequence 1334, Appl		
c	36	35	4.0	696	4	US-09-583-110-1334	Sequence 176, Appl		
c	37	34.6	4.0	789	4	US-09-107-433-176	Sequence 8688, Appl		
c	38	34.6	4.0	921	4	US-09-227-767-8688	Sequence 23970, A		
c	39	34.6	4.0	921	4	US-09-270-767-23970	Sequence 59, Appl		
c	40	34.6	4.0	1086	4	US-09-248-796A-59	Sequence 2564, Appl		
c	41	34.2	3.9	3028	4	US-09-248-796A-2564	Sequence 13498, A		
c	42	34.2	3.9	343352	4	US-09-949-016-13198	Sequence 15605, A		
c	43	34.2	3.9	55806	4	US-09-949-016-13198	Sequence 37936, A		



SOFTWARE: FastSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/792,014  
 FILING DATE: Herewith  
 CLASSIFICATION: 435  
 PRIOR APPLICATION DATA:  
 APPLICATION NUMBER:  
 FILING DATE:  
 ATTORNEY/AGENT INFORMATION:  
 NAME: Billings, Lucy J.  
 REGISTRATION NUMBER: 36,749  
 REFERENCE/DOCKET NUMBER: PF-0206 US  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: 415-855-0555  
 TELEFAX: 415-845-4166  
 TELEX:  
 INFORMATION FOR SEQ ID NO: 2:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 819 base pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 LIBRARY: Consensus  
 CLONE: Consensus  
 US-08-792-014-2

Query Match 5.3%; Score 46.4%; DB 3; Length 819;  
 Best Local Similarity 50.0%; Pred. No. 0; 0 00029;  
 Matches 116; Conservative 0; Mismatches 116; Indels 0; Gaps 0;

Qy 540 ATATCTTCACTTGGAAACAGATTGACTGAATAATGAGCTGAACTGATGCCAGTCCTTCATCA 599  
 Db 546 ATTTCCTGGATGGCAATGAATGACATTAGCTGATTCACCTGCTGCCAACTCTGAT 605

Qy 600 TATTGGAATTATTGGATTGTCATCTCTGGATTGCAATTTCACATCT 659  
 Db 606 TGTCAAGGTGTGGCCAAAAAAATATGCAACTTGTGATTGACTGGCAT 665

Qy 660 CTGGGTTATATCCCTACCTGATCACTGATCAAGGAGTTATTGGAGTGTCCCGCGA 719  
 Db 666 CTGGAGATACTTAACTATGCACTAGTGGACGGTTACCAATACCTGTCAGTGA 725

Qy 720 TCAGGACATTATTCTACATCACTAAAGAACATGAAATGGATCTGTCAAAATCAA 771  
 Db 726 TAAGGGGTTGAAATAGCATATGGATGTAGCCAAAAGACTCACCAAGTAA 777

RESULT 6  
 US-09-443-948-2  
 Sequence 2, Application US/09443948  
 Patent No. 6228616

GENERAL INFORMATION:  
 APPLICANT: Bandman, Olga  
 Colli, Surya K.  
 TITLE OF INVENTION: NOVEL HUMAN ANTON CHANNEL  
 NUMBER OF SEQUENCES: 4  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Incyte Pharmaceuticals, Inc.  
 STREET: 3174 Porter Drive  
 STATE: CA  
 COUNTRY: USA  
 ZIP: 94304

COMPUTER READABLE FORM:  
 MEDIUM TYPE: Diskette  
 COMPUTER: IBM Compatible  
 OPERATING SYSTEM: DOS  
 SOFTWARE: FastSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/09/443,948  
 FILING DATE: 19-No. 6228616-1999  
 CLASSIFICATION: <Unknown>

```

NAME: Billings, Lucy J. ; SEQ ID NO: 815
REGISTRATION NUMBER: 36,749 ; LENGTH: 1652
REFERENCE/DOCKET NUMBER: PP-0206 US ; TYPE: DNA
TELECOMMUNICATION INFORMATION: ; ORGANISM: Homo sapiens
TELEPHONE: 415-855-0555 ; FEATURE:
TELEFAX: 415-845-4166 ; NAME/KEY: CDS
TELEX: <Unknown> ; LOCATION: (232) .. (975)
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 819 base pairs ; US-09-620-312D-815
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: Consensus
CLONE: Consensus
SEQUENCE DESCRIPTION: SEQ ID NO: 2:
us-09-690-196-2

Query Match 5.3%; Score 46.4; DB 4; Length 819;
Best Local Similarity 50.0%; Pred. No. 0.0039;
Matches 116; Conservative 0; Mismatches 116; Indels 0; Gaps 0;
Qy 540 ATATCTACTTGGAAACAGTTGACTGAAATGAAATGACTGAACTGATGCCAGTCCTCATCA 599
Db 546 ATTTCCTGATGCCAATGAAATGAACTGAACTGATGCCAAACTGCTAT 605
Qy 600 TATTGGAATTATTGGATTGCACTCTGGATTGATATTCATCTCATCT 659
Db 606 TGTCAAGGGGGCCAAAAAAATATGCCAACATTGTATTCAGAAATGACTGGCT 665
Qy 660 CTGGGTATATCCTCACTGCATACCGTACAGCAGATTATTGAGAGTTGCTCCGCCGA 719
Db 666 CTGGAGACCTTAACTTACATAGCATAGTAGCTGGACGGTCACTGTCCAGTGA 725
Qy 720 TCAAGGACATTATTCATCACTATAAGAACAAATGAAATCTGTCAAAATCAA 771
Db 726 TAAGGGTTGAAATAGCATATGATGATGAGCTTGTAAAGACTCAGGTAA 777

RESULT 8
US-09-949-016-5780
; Sequence 5780, Application US/0949016
; Patent No. 6812319
; GENERAL INFORMATION:
; APPLICANT: J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIORITY FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/1241,755
; PRIOR PILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-01-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIORITY FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 2070012
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO: 5780
; LENGTH: 1689
; TYPE: DNA
; ORGANISM: Human
; US-09-949-016-5780

Query Match 5.0%; Score 43.4; DB 4; Length 1689;
Best Local Similarity 52.5%; Pred. No. 0.0039;
Matches 95; Conservative 0; Mismatches 86; Indels 0; Gaps 0;
Qy 549 TGGAAACAGTATGACTGAAATGACTGAACTGATGCCACGTTCTCATATTCGAT 608
Db 756 TGGGACCAAGAAATATCGTACATGGACTTCAAGGATTCAGGTCTGGCTTA 815
Qy 609 TATTGGAATTGTCACTCTGGATTGATATTCATCTCATATTCGAT 668
Db 816 TGCTCCAAAGAAATATCGTACATGGACTTCAAGGAGTTCTGGCTTA 875
Qy 669 TATCTCTACTGATGAACTGATGCCAGTATTGAGATTGCTGGCGATCAGGACAT 728
Db 876 TCTCCACAATGCTATGCCGTGAAGAATTACCCACGTCTGAAGACAAGAAAT 935
Qy 729 T 729

RESULT 9
US-09-949-016-5780
; Sequence 5780, Application US/0949016
; Patent No. 6812319
; GENERAL INFORMATION:
; APPLICANT: J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIORITY FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/1241,755
; PRIOR PILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-01-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIORITY FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 2070012
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO: 5780
; LENGTH: 1689
; TYPE: DNA
; ORGANISM: Human
; US-09-949-016-5780

Query Match 5.0%; Score 43.4; DB 4; Length 1689;
Best Local Similarity 52.5%; Pred. No. 0.0039;
Matches 95; Conservative 0; Mismatches 86; Indels 0; Gaps 0;
Qy 549 TGGAAACAGTATGACTGAAATGACTGAACTGATGCCACGTTCTCATATTCGAT 608
Db 746 TGGGACCAAGAAATATCGTACATGGACTTCAAGGATTCAGGTCTGGCTTA 865
Qy 609 TATTGGAATTGTCACTCTGGATTGATATTCATCTCATATTCGAT 668
Db 806 TGCTCCAAAGAAATATCGTACATGGACTTCAAGGAGTTCTGGCTTA 875
Qy 669 TATCTCTACTGATGAACTGATGCCAGTATTGAGATTGCTGGCGATCAGGACAT 728
Db 866 TCTCCACAATGCTATGCCGTGAAGAATTACCCACGTCTGAAGACAAGAAAT 925
Qy 729 T 729

```

Db 926 T 926

RESULT 9

US-09-710-794-3

Sequence 3, Application US/0910794

Patent No. 6570369

GENERAL INFORMATION

APPLICANT: Holloway, James L.

APPLICANT: Gao, Zeren

TITLE OF INVENTION: NOVEL CRIB PROTEIN ZMSB1

FILE REFERENCE: 99-76

CURRENT APPLICATION NUMBER: US/09/710,794

CURRENT FILING DATE: 2000-11-09

PRIOR APPLICATION NUMBER: US 60/164,685

PRIOR FILING DATE: 1999-11-10

NUMBER OF SEQ ID NOS: 31

SEQUENCE: FastSEQ for Windows Version 3.0

SEQ ID NO 3

LENGTH: 1068

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE: INFORMATION: Degenerate polynucleotide sequence for human zmsel

OTHER INFORMATION: misc\_feature

LOCATION: (1) ..(1068)

OTHER INFORMATION: n = A,T,C or G

US-09-710-794-3

Query Match 4.7%; Score 41; DB 4; Length 1068;

Best Local Similarity 25.7%; Pred. No. 0.017;

Matches 76; Conservative 59; Mismatches 160; Indels 1; Gaps 1;

Qy 158 GAGTTGCAAGCTGAAAGTGTCAAGCTGAATTCAGGATTTAAGGAACT 217

Db 187 GARWSNTNGAYGARCCRNWSNWSNWSNARGMWSNTNTNWSNMGNAAR 246

Qy 218 TTCTCGGAGCACACCCGATTATGATGTGAAAGGAAAAAGCTGCAATACGTATA 277

Db 247 TTYYMGNGGNTWSNARGMWSNCAWSNNTNACNMNGGNGARMGNGCARMGNGAYATG 306

Qy 278 ATGGAGAGATGGAGGGATTCCTCAT-TGGCAAAGGAAATTCAATGTTCCACTCTT 336

Db 307 YTNGGNNNTYNTMNGNAYWSNGCNNTNTYTGNAARAYGNCNTGNSNNTCNCARYN 366

Qy 337 GAAAAGGATCCATCGCTGAAAGAGAAACTGTACAGGAACACTCAAACCTGTC 396

Db 367 AYGARARARGCNGNARARGCNAAACNNWSNARNTCCNAARNSNTNWSNNSN 426

Qy 397 CNGTNAARAAACNAAAGTAGAGTTGCTGATAAGGAAAAAAGGCCATCGAGGTGAAGA 452

Db 427 CNGTNAARAAACNAAAGTAGAGTTGCTGATAAGGAAAAAAGGCCATCGAGGTGAAGA 482

RESULT 10

US-09-949-016-107953

Sequence 10, Application US/0949016

Patent No. 6812339

GENERAL INFORMATION

APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

FILE REFERENCE: CLO01307

CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SEQUENCE: FastSEQ for Windows Version 4.0

SEQ ID NO 107954

LENGTH: 601

TYPE: DNA

ORGANISM: Human

US-09-949-016-107954

Query Match 4.6%; Score 39.8; DB 4; Length 601;

Best Local Similarity 52.0%; Pred. No. 0.03;

Matches 89; Conservative 0; Mismatches 82; Indels 0; Gaps 0;

Qy 324 GTTCCACTCTGAAAGGTCCATCGGTGAGGAGAACTGTACAGGAA 383

Db 367 TGTAAACTCTGGAAAGTAACTTAAAGGAAAGACAGTAACAGAA 426

Qy 384 CTTCAGATCTGTTCTGGAGAAAGTAGAGCTTCAATGAGCTGTGAG 443

Db 427 ATACTCTATGTCCTGTGATAAGGACTAAATCTAGGAGCTCTCA 486

Qy 444 AGTTGAAAGATCTCAGCACAGATAAAAGTICACTACAATGAGCTGTGAG 494

Db 488 AGTTATAACTCTGGAAAGTAACTTAAAGGAAAGACAGTAACAGAA 537

RESULT 11

US-09-949-016-107954

Sequence 107954, Application US/0949016

Patent No. 6812339

GENERAL INFORMATION

APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

FILE REFERENCE: CLO01307

CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SEQUENCE: FastSEQ for Windows Version 4.0

SEQ ID NO 107954

LENGTH: 601

TYPE: DNA

ORGANISM: Human

US-09-949-016-107954

Query Match 4.6%; Score 39.8; DB 4; Length 601;

Best Local Similarity 52.0%; Pred. No. 0.03;

Matches 89; Conservative 0; Mismatches 82; Indels 0; Gaps 0;

Qy 324 GTTCCACTCTGAAAGGTCCATCGGTGAGGAGAACTGTACAGGAA 383

Db 368 TGTAAACTCTGGAAAGTAACTTAAAGGAAAGACAGTAACAGAA 427

Qy 384 CTTCAGATCTGTTCTGGAGAAAGTAGAGCTTCAATGAGCTGTGAG 443

Db 428 ATACTCTATGTCCTGTGATAAGGACTAAATCTAGGAGCTCTCA 487

Qy 444 AGTTGAAAGATCTCAGCACAGATAAAAGTICACTACAATGAGCTGTGAG 494

Db 488 AGTTATAACTCTGGAAAGTAACTTAAAGGAAAGACAGTAACAGAA 538

RESULT 12

US-09-949-016-14720/C

Sequence 14720, Application US/0949016

Patent No. 6812339

GENERAL INFORMATION

APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

FILE REFERENCE: CLO01307

CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF  
FILE REFERENCE: CL001307  
CURRENT APPLICATION NUMBER: US/09/949,016  
CURRENT FILING DATE: 2000-04-14  
PRIORITY APPLICATION NUMBER: 60/241,755  
PRIORITY FILING DATE: 2000-10-20  
PRIORITY APPLICATION NUMBER: 60/237,768  
PRIORITY FILING DATE: 2000-10-03  
PRIORITY APPLICATION NUMBER: 60/231,498  
PRIORITY FILING DATE: 2000-09-08  
NUMBER OF SEQ ID NOS: 207012  
SOFTWARE: FastSEQ for Windows Version 4.0  
SEQ ID NO: 14720  
LENGTH: 390830  
TYPE: DNA  
ORGANISM: Human  
FEATURE: misc\_feature  
NAME/KEY: misc\_feature  
LOCATION: (1)..(390830)  
OTHER INFORMATION: n = A,T,C or G

US-09-949-016-14720

Query Match 4.6%; Score 39.8; DB 4; Length 390890;  
Best Local Similarity 52.0%; Pred. No. 1..2;  
Matches 89; Conservative 0; Mismatches 82; Indels 0; Gaps 0;

Qy 324 TGTCGCACTCTTGAARAGGATCCATCGCTGAGAAGAACTTGTACAGAA 383  
Db 314235 TGTAAACCTCTGGAAAGATACTTAAAGAGAAACAGAAAGAA 314176

Qy 384 CTTCAAACTGTCTCGGACAAAAGTAGTGGTTCTGATAAGGAAAAGGCCATCGAG 443  
Db 314175 ATACTATGTCTCTGAAATAAGACTAACTCTAGGTATAAAATTCTGAGACAGCTCA 314116

Qy 444 AGCTGAGATCTCCACACAGATTAAGGTCACATCATCGAGCTGTGA 494  
Db 314115 AGTTATAACTCCAGTCCACATATGATGTTCCATCTGTGCTTGTGA 314065

RESULT 13  
US-09-533-029-49  
; Sequence 4.9, Application US/09533029  
; Patent No. 6664446  
; GENERAL INFORMATION:  
; APPLICANT: Heard, Jacqueline  
; APPLICANT: Brou, Pierre  
; APPLICANT: Riechmann, Jose-Luis  
; APPLICANT: Keddie, James  
; APPLICANT: Pineda, Omaira  
; APPLICANT: Adam, Luc  
; APPLICANT: Samaha, Raymond  
; APPLICANT: Zhang, James  
; APPLICANT: Yu, Guo-Liang  
; APPLICANT: Ratcliffe, Oliver  
; APPLICANT: Pilgrim, Marsha  
; APPLICANT: Jiang, Cai-Zhong  
; APPLICANT: Reuber, Lynne  
; TITLE OF INVENTION: DISEASE-INDUCED POLYNUCLEOTIDES  
; FILE REFERENCE: MBL-010  
; CURRENT APPLICATION NUMBER: US/09/533,029  
; EARLIER APPLICATION NUMBER: 2000-03-22  
; EARLIER FILING DATE: 1999-03-23  
; NUMBER OF SEQ ID NOS: 121  
; SOFTWARE: Patentin Ver. 2.1  
; SEQ ID NO: 49  
; LENGTH: 1529  
; TYPE: DNA  
; ORGANISM: Arabidopsis thaliana  
; FEATURE: G553  
; OTHER INFORMATION: G553  
US-09-533-029-49

Query Match 4.5%; Score 39.2; DB 4; Length 1529;  
Best Local Similarity 58.6%; Pred. No. 0..08; Mismatches 48; Indels 0; Gaps 0;  
Matches 68; Conservative 0;

Qy 150 TGAGATTGGACTTGGACGGACTGCAACGTGAACTCTGAAGCTTAA 209  
Db 249 TGAGGTGATGCTGACCCAGGCGATGATAACATAGTGTAACTCTGTAA 308

Qy 210 GAAGAACTTCTCGGACAAACCCGATTGTTAGAGGABAAGAGCTA 265  
Db 309 TAATAACTCTTGAGCAGAACCTCGACTAATATGTCAGGGAACCGGA 364

RESULT 14  
US-09-806-708B-22  
; Sequence 22, Application US/09806708B  
; Patent No. 6784312  
; GENERAL INFORMATION:  
; APPLICANT: The University of British Columbia  
; TITLE OF INVENTION: Regulation of Embryonic Transcription in Plants  
; FILE REFERENCE: 4810-58741  
; CURRENT APPLICATION NUMBER: US/09/806,708B  
; CURRENT FILING DATE: 2001-04-03  
; PRIORITY APPLICATION NUMBER: US 60/147,133  
; PRIORITY FILING DATE: 1999-08-04  
; SOFTWARE: Patentin version 3.0  
; SEQ ID NO: 23  
; LENGTH: 1141  
; TYPE: DNA  
; ORGANISM: Artificial sequence  
; FEATURE:  
; NAME/KEY: promoter  
; LOCATION: (1)..(1141)  
; OTHER INFORMATION: consensus sequence of A.t., L.a., and B.n. FAEB1 promoter  
US-09-806-708B-22

Query Match 4.4%; Score 38.4; DB 4; Length 1141;  
Best Local Similarity 8.5%; Pred. No. 0..12;  
Matches 36; Conservative 183; Mismatches 207; Indels 0; Gaps 0;

Qy 284 AGATTTGAAAGGACGGATCTTCATTTGGAAAGGAACTTCAATGTTGCGAG 403  
Db 176 DDDTGYHMMNNNGCETVIMVYKTDWDMSBKRNMNGBMWKNWSDVITYWWDDMCK 295

Qy 344 ATCCATTCGGTGTAGAGAATAGGAACTTCAACCTGTTCCGTGAG 355  
Db 236 RKVREWVWRGRMENYMWABTAIRRYNGWBTAMAYERWTHNNNNNAKAMCRAKWY 295

Qy 404 CAAAGTAGTGGATCTGCTATAGGAAAAGGCCATCGAGAGTTGAAAGATCTCCAGCAC 463  
Db 296 GWNRLBVNSTCTTWSKTTKVRSTCWNCRAGDKDEIKWWKWSAAMGVYWNNNNNNN 355

Qy 464 AGATTAAGTCACTACAATCGTGTGAGAACATTCGAATTTGATCAGTGTCTAT 523  
Db 356 TYKKRHBARWDWTHWSANKWTHANAHYSRKWPTBYKRKTMVNNNGTMMWRMWAHY 415

Qy 524 CCGACAGAAATCTGATATCTACTGGAAACAGTATGACTGTAATGACTGTAAGTGA 583  
Db 416 RMDMMWBGTYNNNNNGRTTYGWTRKRNKRWWTYKWRANNCKWRADWKTCRHNNTWWQM 475

Qy 584 TGCCCACTGCTCTCATATATCGAATATTGGATGTCATCTCTGATCATTCAC 643  
Db 476 KTYNNCYWKSMTNKSHEBAAYTWWWWWWRYAHANNNNNWDYWWKACTWVXYBVCSK 535

Qy 644 ATAATTTCATCTCATCTCTGGCTTATATCCTCACTGTCAGAGCATTATG 703  
Db 536 WWWWYAAWYTKSSNNWTSYYRWKTNNSWRSDTRSMGRANNYARAHGYKWNTRWNB 595

Qy 704 AGAGTT 709  
Db 596 WSHTWB 601

RESULT 15  
US-09-949-016-12656 Application US/09349016  
| Sequence 12656 , Application US/09349016  
| Patent No. 6812339  
| GENERAL INFORMATION  
| APPLICANT: VENTER, J. Craig et al.  
| TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED  
| WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF  
| FILE REFERENCE: CL001307  
| CURRENT APPLICATION NUMBER: US/09/949,016  
| CURRENT FILING DATE: 2000-04-14  
| PRIOR APPLICATION NUMBER: 60/1241,755  
| PRIOR FILING DATE: 2000-10-20  
| PRIOR APPLICATION NUMBER: 60/237,768  
| PRIOR FILING DATE: 2000-10-03  
| PRIOR APPLICATION NUMBER: 60/231,498  
| PRIOR FILING DATE: 2000-09-08  
| NUMBER OF SEQ ID NOS: 207012  
| SOFTWARE: FastSEQ for Windows Version 4.0  
| SEQ ID NO: 12656  
| LENGTH: 253345  
| TYPE: DNA  
| ORGANISM: Human  
| US-09-949-016-12656

Query Match 4.4%; Score 38.4; DB 4; Length 253345;  
Best Local Similarity 57.5%; Pred. No. 2,6;  
Matches 69; Conservative 0; Mismatches 51; Indels 0; Gaps 0;  
Qy 438 ATCGAGAGTTGAAAGATCTCCAGCACAGATTAAAGTTCACTACAATCGAGCTGTGAGCA 497  
Db 17149 ATCATCATGCTATAATTATGCAAAAGATTAAAGTTAAAGTCATGAAACTCCCTATGAAA 17208  
Qy 498 ACTATCCGATATTGATGAGTTGCTATCGAGGAAATCTCGATATCTACTTGGAAACAG 557  
Db 17209 CACTTCCGATCAAGACCTACAGCAAGCTGATAAAATTAACATTAAACATGAAAATG 17268

Search completed: March 10, 2005, 02:24:01  
Job time : 202 secs

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AT TACOMA MINE  
Maximum Match 100%  
Minimum Match 45%  
Fictive Factor 4.5 Gammma = 4.00

Database : N\_Geneseq\_16Dec04 : \*  
 1: geneseqn19806; \*  
 2: geneseqn19908; \*  
 3: geneseqn20008; \*  
 4: geneseqn2001as; \*  
 5: geneseqn2001bs; \*  
 6: geneseqn2002as; \*  
 7: geneseqn2002bs; \*  
 8: geneseqn2003as; \*  
 9: geneseqn2003bs; \*  
 10: geneseqn2003cs; \*  
 11: geneseqn2003ds; \*  
 12: geneseqn2004as; \*  
 13: geneseqn2004bs; \*  
 RESULT 1  
 ADS96405  
 ID ADS96405 standard; cDNA; 783 BP.  
 XX  
 AC  
 ADS96405;  
 XX  
 DT 02-DEC-2004 (first entry)  
 XX  
 DE Drosophila melanogaster protein coding sequence, SEQ ID 26.  
 XX  
 KW Insecticide; Antiparasitic; Antihelminthic; gene; ds.  
 XX  
 OS Drosophila melanogaster.

Required. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

result No.	Score	Query	Match	Length	DB	ID	Description
1	76.4	8.8	783	13	AD596405		Ad596405 Drosophil
2	76.4	8.8	1063	4	AB17133		Ab17133 Drosophil
3	63	7.2	786	3	AAAC5377		Aac5377 Cat flea
4	63	7.2	786	3	AAAC5376		Aac5376 Cat flea
5	63	7.2	2383	3	AAAC5375		Aac5375 Cat flea
6	63	7.2	2383	3	AAAC5374		Aac5374 Cat flea
7	51.2	5.9	2000	8	ADA1938		Ada1938 Rice Gene
8	48	5.5	2238	11	AD02307		Ad02307 Human cDN
9	48	5.5	2887	9	ADA4507		Ada4507 Human cDN
10	48	5.5	4290	8	ACCC3320		Acc3320 Chloride
11	48	5.5	4318	4	AAK15103		Aak15103 Human pol
12	48	5.5	4318	4	AAI57882		Aai57882 Human pol
13	48	5.5	4318	8	ACF12844		Acf12844 Human cer
14	48	5.5	4318	13	AD225580		Adr225580 Breast ca
15	48	5.5	4318	13	AD23232		Adp23232 PRO polyp
16	48	5.5	4707	11	AD86762		Adm86762 Human cDN
17	46.4	5.3	819	2	AAV3260		Aav3260 DNA encod
18	46.4	5.3	819	10	AD56392		Adj56392 Human cDN
19	46	5.3	4357	4	AAK2487		Aak2487 Human pol
20	46	5.3	4357	4	AAAT9668		Aat9668 Human pol

PS XX The present invention relates to a method for identifying a compound that  
CC inhibits the activity of a protein essential for *Drosophila* viability.  
CC The method comprises: (a) expressing in a recombinant host a DNA sequence  
CC encoding a protein essential for *Drosophila*; (b) testing  
CC compounds suspected of having the ability to inhibit the activity of the  
CC protein expressed in (a); and identifying a compound tested in (b) that  
CC inhibits the activity of the protein. The method is useful in identifying no  
CC

PS Claim 1; SEQ ID NO 26; 5/pp; English.





Db 519 GATGCCAGGTTACACACATCAGGTGGCCAGTATTCTGCAATTGAAATCC 578  
 Qy 642 ACATATTCTACTCATCTGGCTTATATCCTCATGGATAACCGTACAGGATTAT 701  
 Db 579 GAGCAATCAACCGCTTATGGCTTATGTTACATGTTACAGGTGACGGATTCC 638  
 Db 702 TGAGGTTGTCGGCGATAGGATTTCATCACTATAAGACAGATGCTGT 761  
 Qy 639 CCAGTGTCGCCAGCGCAAGATATCAGATACCACTATAACTGCAAGGAGAT 698  
 Db 762 CACAAATCAC 772  
 Db 699 CAGCAATAACC 709

RESULT 5

AAC95375/c  
 ID AAC95375 standard; cDNA; 2383 BP.  
 XX  
 AC AAC95375;  
 XX  
 DT 19-FEB-2001 (first entry)  
 XX Cat flea HMT CL intracellular channel cDNA complement, SEQ ID NO:1874.  
 XX Cat flea; hindgut and Malpighian tubule nucleic acid; HMT;  
 XX file infestation; vaccine; antiparasitic; therapeutic target; diagnosis;  
 KW detection; ss.  
 XX Ctenocephalides felis.  
 OS WO20061621-A2.  
 XX  
 PD 19-OCT-2000.  
 XX  
 PF 07-APR-2000; 2000WO-US009437.  
 XX  
 PR 09-APR-1999; 99US-0128704P.  
 XX  
 PA (HESKA) HESKA CORP.  
 XX  
 Brandt KS, Gaines PJ, Stinchcomb DT, Wisniewski N;  
 XX  
 WPI; 2000-656323/63.  
 DR P-PDB; AAB29622.

XX Flea Malpighian tubule and nerve cord tissue derived nucleic  
 PT acids useful for the prevention, diagnosis and treatment of flea  
 PT infestations.  
 XX  
 PS Claim 1; Page 900-901; 964pp; English.

XX The invention relates to novel cat flea (*Ctenocephalides felis*) nucleic  
 CC acids which are expressed in hindgut and Malpighian tubule (HMT) tissue  
 CC or head and nerve cord (HNC) tissue. The invention also relates to the  
 CC encoded proteins. The invention additionally encompasses expression of  
 CC constructs, recombinant viruses and recombinant cells comprising the  
 CC nucleic acids of the invention, recombinant production of the proteins,  
 CC antibodies against the proteins, a method of identifying inhibitors of  
 CC the proteins, and compositions comprising the inhibitors for  
 CC administration to an animal. The nucleic acids, and the proteins they  
 CC encode may be used in the prevention, treatment and diagnosis of diseases  
 CC associated with flea infestations. For example, the nucleic acids may be  
 CC used to produce an HMT or HNC protein according to standard recombinant  
 CC DNA methodology by inserting the nucleic acids into a host cell and  
 CC culturing the cell to express the protein. The HMT and HNC nucleic acids  
 CC may also be used as DNA probes in diagnostic assays (e.g., PCR) to detect  
 CC and quantitate the presence of cat flea or other homologous nucleic acid  
 CC sequences in samples. They may also be used to study the expression and  
 CC function of the proteins and their role in metabolism. The HMT and HNC  
 CC proteins may be used as antigens in the production of specific  
 CC antibodies, and in assays to identify modulators (agonists and  
 CC antagonists) of HMT and/or HNC protein expression and activity. The anti-

CC HMT/HNC protein antibodies and antagonists may also be used to  
 CC downregulate protein expression and activity. The antibodies may also be  
 CC used as diagnostic agents for detecting the presence of flea polyptides  
 CC in samples (e.g., by enzyme linked immunosorbent assay (ELISA)). The  
 CC present sequence represents a cat flea HMT cDNA of the invention  
 XX

Sequence 2383 BP; 812 A; 374 C; 354 G; 835 T; 0 U; 8 Other;  
 Query Match 7.2%; Score 63; DB 3; Length 2383;  
 Best Local Similarity 50.2%; Pred. No. 5.3e-08;  
 Matches 156; Conservative 0; Mismatches 155; Indels 0; Gaps 0;

Qy 462 ACAGATTAAGTCTACAAATCGAGTGTGAGCAATATGCAATTGATCAGTCT 521  
 Db 1926 AACGATGACGTAAACGATCTGAACTCTGAGCCCTGAGAAATCACACACATT 1867  
 Qy 522 ATCCGAGAGAAAATCTCGATATCTACTGAAATGACTGAATATGACTGTGAACT 581  
 Db 1866 GGCGGGCGGGAGAGATTCTAACGGGACACCCGTCGTCGAGGAACT 1807  
 Qy 582 GAGCCAGGTCTCTCTCATATTGCAATTGATGATGATGATGATGATGATATCC 641  
 Db 1806 GATGCCAGGTTACACATCAGGTCGCCCAAGTATTCTGCTGATTGAAATCC 1747  
 Qy 642 ACATAATTCTACTCATCTGCGCTATATCCTCATCTGATACCGAGCATTT 701  
 Db 1746 GAGCAATTAACGGCTTATGSGTTATGATCATGATGATGATGATGATGAT 1687  
 Qy 702 TGAGAGTTGTCGGCGATCAGGACATTATCATACTATAAGAACAAATGATTGTT 761  
 Db 1686 CCAGTCGTGCCAGCGACCAAGATCATACCACTATAACTGCAACGAGGAT 1627  
 Qy 762 CACAAATCAAC 772  
 Db 1626 CAGCAATAACC 1616

RESULT 6

AAC95374  
 ID AAC95374 standard; cDNA; 2383 BP.  
 XX  
 AC AAC95374;  
 XX  
 DT 19-FEB-2001 (first entry)  
 XX Cat flea HMT C1 intracellular channel cDNA, SEQ ID NO:1872.  
 XX  
 DE Cat flea hindgut and Malpighian tubule nucleic acid; HMT;  
 KW flea infestation; vaccine; antiparasitic; therapeutic target; diagnosis;  
 KW detection; ss.  
 XX  
 Ctenocephalides felis.  
 OS  
 XX  
 WO20061621-A2.  
 XX  
 PR 09-APR-1999; 99US-0128704P.  
 XX  
 PA (HESKA) HESKA CORP.  
 XX  
 Brandt KS, Gaines PJ, Stinchcomb DT, Wisniewski N;  
 XX  
 WPI; 2000-656323/63.  
 DR P-PDB; AAB29622.

XX  
 PS Claim 1; Page 900-901; 964pp; English.

XX The invention relates to novel cat flea (*Ctenocephalides felis*) nucleic  
 CC acids which are expressed in hindgut and Malpighian tubule (HMT) tissue  
 CC or head and nerve cord (HNC) tissue. The invention also relates to the  
 CC encoded proteins. The invention additionally encompasses expression of  
 CC constructs, recombinant viruses and recombinant cells comprising the  
 CC nucleic acids of the invention, recombinant production of the proteins,  
 CC antibodies against the proteins, a method of identifying inhibitors of  
 CC the proteins, and compositions comprising the inhibitors for  
 CC administration to an animal. The nucleic acids, and the proteins they  
 CC encode may be used in the prevention, treatment and diagnosis of diseases  
 CC associated with flea infestations. For example, the nucleic acids may be  
 CC used to produce an HMT or HNC protein according to standard recombinant  
 CC DNA methodology by inserting the nucleic acids into a host cell and  
 CC culturing the cell to express the protein. The HMT and HNC nucleic acids  
 CC may also be used as DNA probes in diagnostic assays (e.g., PCR) to detect  
 CC and quantitate the presence of cat flea or other homologous nucleic acid  
 CC sequences in samples. They may also be used to study the expression and  
 CC function of the proteins and their role in metabolism. The HMT and HNC  
 CC proteins may be used as antigens in the production of specific  
 CC antibodies, and in assays to identify modulators (agonists and  
 CC antagonists) of HMT and/or HNC protein expression and activity. The anti-

CC HMT/HNC protein antibodies and antagonists may also be used to  
 CC downregulate protein expression and activity. The antibodies may also be  
 CC used as diagnostic agents for detecting the presence of flea polyptides  
 CC in samples (e.g., by enzyme linked immunosorbent assay (ELISA)). The  
 CC present sequence represents a cat flea HMT cDNA of the invention  
 XX

Sequence 2383 BP; 812 A; 374 C; 354 G; 835 T; 0 U; 8 Other;  
 Query Match 7.2%; Score 63; DB 3; Length 2383;  
 Best Local Similarity 50.2%; Pred. No. 5.3e-08;  
 Matches 156; Conservative 0; Mismatches 155; Indels 0; Gaps 0;

Qy 462 ACAGATTAAGTCTACAAATCGAGTGTGAGCAATATGCAATTGATCAGTCT 521  
 Db 1926 AACGATGACGTAAACGATCTGAACTCTGAGCCCTGAGAAATCACACACATT 1867  
 Qy 522 ATCCGAGAGAAAATCTCGATATCTACTGAAATGACTGAATATGACTGTGAACT 581  
 Db 1866 GGCGGGCGGGAGAGATTCTAACGGGACACCCGTCGTCGAGGAACT 1807  
 Qy 582 GAGCCAGGTCTCTCTCATATTGCAATTGATGATGATGATGATGATGATATCC 641  
 Db 1806 GATGCCAGGTTACACATCAGGTCGCCCAAGTATTCTGCTGATTGAAATCC 1747  
 Qy 642 ACATAATTCTACTCATCTGCGCTATATCCTCATCTGATACCGAGCATTT 701  
 Db 1746 GAGCAATTAACGGCTTATGSGTTATGATCATGATGATGATGATGAT 1687  
 Qy 702 TGAGAGTTGTCGGCGATCAGGACATTATCATACTATAAGAACAAATGATTGTT 761  
 Db 1686 CCAGTCGTGCCAGCGACCAAGATCATACCACTATAACTGCAACGAGGAT 1627  
 Qy 762 CACAAATCAAC 772  
 Db 1626 CAGCAATAACC 1616

RESULT 7

AAC95374  
 ID AAC95374 standard; cDNA; 2383 BP.  
 XX  
 AC AAC95374;  
 XX  
 DT 19-FEB-2001 (first entry)  
 XX Cat flea HMT C1 intracellular channel cDNA, SEQ ID NO:1872.  
 XX  
 DE Cat flea hindgut and Malpighian tubule nucleic acid; HMT;  
 KW flea infestation; vaccine; antiparasitic; therapeutic target; diagnosis;  
 KW detection; ss.  
 XX  
 Ctenocephalides felis.  
 OS  
 XX  
 WO20061621-A2.  
 XX  
 PR 09-APR-1999; 99US-0128704P.  
 XX  
 PA (HESKA) HESKA CORP.  
 XX  
 Brandt KS, Gaines PJ, Stinchcomb DT, Wisniewski N;  
 XX  
 WPI; 2000-656323/63.  
 DR P-PDB; AAB29622.

XX  
 PS Claim 1; Page 900-901; 964pp; English.

XX The invention relates to novel cat flea (*Ctenocephalides felis*) nucleic  
 CC acids which are expressed in hindgut and Malpighian tubule (HMT) tissue  
 CC or head and nerve cord (HNC) tissue. The invention also relates to the  
 CC encoded proteins. The invention additionally encompasses expression of  
 CC constructs, recombinant viruses and recombinant cells comprising the  
 CC nucleic acids of the invention, recombinant production of the proteins,  
 CC antibodies against the proteins, a method of identifying inhibitors of  
 CC the proteins, and compositions comprising the inhibitors for  
 CC administration to an animal. The nucleic acids, and the proteins they  
 CC encode may be used in the prevention, treatment and diagnosis of diseases  
 CC associated with flea infestations. For example, the nucleic acids may be  
 CC used to produce an HMT or HNC protein according to standard recombinant  
 CC DNA methodology by inserting the nucleic acids into a host cell and  
 CC culturing the cell to express the protein. The HMT and HNC nucleic acids  
 CC may also be used as DNA probes in diagnostic assays (e.g., PCR) to detect  
 CC and quantitate the presence of cat flea or other homologous nucleic acid  
 CC sequences in samples. They may also be used to study the expression and  
 CC function of the proteins and their role in metabolism. The HMT and HNC  
 CC proteins may be used as antigens in the production of specific  
 CC antibodies, and in assays to identify modulators (agonists and  
 CC antagonists) of HMT and/or HNC protein expression and activity. The anti-

CC HMT/HNC protein antibodies and antagonists may also be used to  
 CC downregulate protein expression and activity. The antibodies may also be  
 CC used as diagnostic agents for detecting the presence of flea polyptides  
 CC in samples (e.g., by enzyme linked immunosorbent assay (ELISA)). The  
 CC present sequence represents a cat flea HMT cDNA of the invention  
 XX

Sequence 2383 BP; 812 A; 374 C; 354 G; 835 T; 0 U; 8 Other;  
 Query Match 7.2%; Score 63; DB 3; Length 2383;  
 Best Local Similarity 50.2%; Pred. No. 5.3e-08;  
 Matches 156; Conservative 0; Mismatches 155; Indels 0; Gaps 0;

Qy 462 ACAGATTAAGTCTACAAATCGAGTGTGAGCAATATGCAATTGATCAGTCT 521  
 Db 1926 AACGATGACGTAAACGATCTGAACTCTGAGCCCTGAGAAATCACACACATT 1867  
 Qy 522 ATCCGAGAGAAAATCTCGATATCTACTGAAATGACTGAATATGACTGTGAACT 581  
 Db 1866 GGCGGGCGGGAGAGATTCTAACGGGACACCCGTCGTCGAGGAACT 1807  
 Qy 582 GAGCCAGGTCTCTCTCATATTGCAATTGATGATGATGATGATGATGATATCC 641  
 Db 1806 GATGCCAGGTTACACATCAGGTCGCCCAAGTATTCTGCTGATTGAAATCC 1747  
 Qy 642 ACATAATTCTACTCATCTGCGCTATATCCTCATCTGATACCGAGCATTT 701  
 Db 1746 GAGCAATTAACGGCTTATGSGTTATGATCATGATGATGATGAT 1687  
 Qy 702 TGAGAGTTGTCGGCGATCAGGACATTATCATACTATAAGAACAAATGATTGTT 761  
 Db 1686 CCAGTCGTGCCAGCGACCAAGATCATACCACTATAACTGCAACGAGGAT 1627  
 Qy 762 CACAAATCAAC 772  
 Db 1626 CAGCAATAACC 1616

RESULT 8

AAC95374  
 ID AAC95374 standard; cDNA; 2383 BP.  
 XX  
 AC AAC95374;  
 XX  
 DT 19-FEB-2001 (first entry)  
 XX Cat flea HMT C1 intracellular channel cDNA, SEQ ID NO:1872.  
 XX  
 DE Cat flea hindgut and Malpighian tubule nucleic acid; HMT;  
 KW flea infestation; vaccine; antiparasitic; therapeutic target; diagnosis;  
 KW detection; ss.  
 XX  
 Ctenocephalides felis.  
 OS  
 XX  
 WO20061621-A2.  
 XX  
 PR 09-APR-1999; 99US-0128704P.  
 XX  
 PA (HESKA) HESKA CORP.  
 XX  
 Brandt KS, Gaines PJ, Stinchcomb DT, Wisniewski N;  
 XX  
 WPI; 2000-656323/63.  
 DR P-PDB; AAB29622.

XX  
 PS Claim 1; Page 896-898; 964pp; English.

The invention relates to novel cat flea (*Ctenocephalides felis*) nucleic acids which are expressed in hindgut and Malpighian tubule (HMT) tissue or head and nerve cord (HNC) tissue. The invention also relates to the encoded proteins. The invention additionally encompasses expression constructs, recombinant viruses and recombinant cells comprising the nucleic acids of the invention, recombinant production of the proteins, antibodies against the proteins, method of identifying inhibitors of the proteins, and compositions comprising the inhibitors for administration to an animal. The nucleic acids, and the proteins they encode may be used in the prevention, treatment, and diagnosis of diseases associated with flea infestations. For example, the nucleic acids may be used to produce an HMT or HNC protein according to standard recombinant DNA methodology by inserting the nucleic acids into a host cell and culturing the cell to express the protein. The HMT and HNC nucleic acids may also be used as DNA probes in diagnostic assays (e.g., PCR) to detect and quantitate the presence of cat flea or other homologous nucleic acid sequences in samples. They may also be used to study the expression and function of the proteins and their role in metabolism. The HMT and HNC proteins may be used as antigens in the production of specific antibodies, and in assays to identify modulators (agonists and antagonists) of HMT and/or HNC protein expression and activity. The anti-HMT/HNC protein antibodies and antagonists may also be used to downregulate protein expression and activity. The antibodies may also be used as diagnostic agents for detecting the presence of flea polypeptides in samples (e.g., by enzyme linked immunosorbent assay (ELISA)). The

CC present sequence represents a cat flea HMT-CDNA of the invention  
XX  
SQ Sequence 2383 BP; 835 A; 354 C; 374 G; 812 T; 0 U; 8 Other;  
Query Match 7 2.2%; Score 63; DB 3; Length 2383;  
Best Local Similarity 50.2%; Pred. No. 5.5e-08;  
Matches 156; Conservative 0; Mismatches 155; Indels 0; Gaps 0;  
OY 462 ACAGTAAAGCTCACTACAACTGGCTTCAGCAAGAAATCCAAATGATCAGGCT 521

the incompatible interaction of plant gene expression relative to expression of the gene in an uninfected plant, in a mutant plant that does not express a gene associated with response to pathogenic infection, or in a corresponding incompatible or compatible interaction. (M1) is useful for conferring resistance to resistance or tolerance to a plant to bacterial, fungal or viral infection. The present sequence was used to illustrate the invention.

Query	Match	Score	Length	DB	8;	Length	2000;
Best Local Similarity	5.9%	Score 51.2;	DB 8;	Length 2000;			
Matches	10.4%	Pred. No. 0.	0.000.9				
Matches	294;	Mismatches	302;	Indels	4;	Gaps	1
Matches	70;	Conservative	294;				
Qy	75	GTCNGGAAATTGATGCTGCGCCGGATTTGGGGCGATTTCTGCTGAGAAATTCTGGATGCGA	134				
Db	1030	GTGTGTAGGTGCTKMRRTYTSMSMTYAMMKCYTMTAYSTSTWKMVWYKMRVAYWSRS	971				
Qy	135	GTTGATGCTCCTTATGATGAGTGGAGTGTGACGAGTGAAGACTGTAACTGTAACTGTGAA	194				
Db	970	RRTWCTGTGRRMATTGCTKRMWAGRNRWMAWCWYCCMWWKTRMTCMWWKTRWTRWTCWY	911				
Qy	195	TCTGAAAGCATTAAAGA----GAACCTTCTCGGAGGACAACCCGATTATGATGAGG	250				
Db	910	TWMMGANYAYAMRRRWRWYKWSWRRMWTMKTWMMKWTWMTCMCKWYMATGWMMWW	851				
Qy	251	AGGAAGAAAGCTGATACACTGATAATCGAGAATTGAGCTGAGGATCTGTGAGGATCTG	310				
Db	850	RYTMYTTCYAMTCAKCKYKMANMKWTTWACWRAWSWRWRAAGMGRWKRKYMKRKYWR	791				
Qy	311	CAAAGGAAATTCAATGTTCCACCTCTTGAAAAGGATCCATCCCTGAGAAGAGAAATAGAGA	370				
Db	790	WRCWKGWARMKMSRTRWKWQKYATRYWKRWQWAMTWMMWSRWRWKSYSRMMWSGRMWSA	731				
Qy	371	ACTTGTCAGGAACTTCAAACTGTTCTGGAGCAAAGTAGACTTCGATAAGGGAAAAA	430				
Db	730	WRYCSRKMKCAKTKYASSARWTKRAKDSYRYYRRTWYKRGKTYTYRYYWRSCRMTRMSK	671				
Qy	431	AGGAGCCATCGAGGAACTTCGAGATCTTCAGCAAGAGTAAACTTCGATAAGGGAAAAA	490				
Db	670	RRKGWAGMSMKCSRMWFGARSHWYSKYSCKSCAKCCTRYTMSSYSTMGMYSKYSKMS	611				
Qy	491	GTGAGGCAACTTCAATATTGATGATGTTGCTGATCCGGAGAAAATCTCGATATCTACTTG	550				
Db	610	WTSMKSYMGKMTCTMITSMKGKSTRRSRNGRSGMSRNMYRMWKKRKRKYMWKWKTCTW	551				
Qy	551	GAACAGTATGACTGAATATGACTGAACTGATGCCACGTCTTCACTCATATTGGAATTA	610				

Db 550 RRCMCYRWGYTMYTTSRMMYTGKARYTTSKRRMWWKPKYCTYYGMMKCSYMM 491  
 Qy 611 TTGGATTGTCACTTCTTGATTCAGATATTCACATCTCGGGCTTATA 670  
 Db 490 RGYCKRACKRCCYAMCWRKAATSGMMWYRKYSKWNRMSTKXMSWYKCRSMKYGK 431  
 Qy 671 TCTCTACTGCTATACCGCTACAGCATTATGAGATGTTGAGCTGTCAGGACATTA 730  
 Db 430 CYGCKWYTCGKTYCYSKYSRVCYKMRMTYKGMNMWYSAVSSMMTWYXX 371  
 Qy 731 TCTCATCTA 740  
 Db 370 AKYWKWYKR 361

RESULT 8  
 ADM02307 standard; cDNA; 2238 BP.  
 XX  
 AC ADM02307,  
 AC ADM02307,  
 XX DT 20-MAY-2004 (first entry)  
 XX DB Human cDNA of the invention SEQ ID NO:992.  
 XX SS; gene; human; gene therapy; diagnostic marker; pharmaceutical.  
 XX KW  
 XX OS Homo sapiens.  
 PN EP1347046-A1.  
 XX PD 24-SEP-2003.  
 XX PF 12-APR-2002; 2002EP-00008400.  
 XX PR 22-MAR-2002; 2002JP-00137785.  
 PA (REAS-) RES ASSOC BIOTECHNOLOGY.  
 XX Isogai T, Sugiyama T, Otsuki T, Wakamatsu A, Sato H, Ishii S;  
 PI Yamamoto J, Isono Y, Hio Y, Otsuka K, Nagai K, Irie R, Tamechika I;  
 PI Seki N, Yoshihawa T, Otsuka M, Nagahari K, Masuho Y;  
 XX WPI: 2003-723558/69.  
 DR P-PSDB; ADM047590.  
 XX New polynucleotides and polypeptides are useful in gene therapy, for  
 PT developing a diagnostic marker or medicines for regulating their  
 PT expression and activity, or as a target of gene therapy.  
 XX PS Claim 1, SEQ ID NO 992; 305pp; English.  
 XX The invention relates to a novel human polynucleotide and the encoded  
 CC polypeptide. A polynucleotide of the invention may have a use in gene  
 CC therapy. An oligonucleotide of the invention ADM06202-ADM06773 is useful  
 CC as a primer for synthesizing the polynucleotide or as a probe for  
 CC detecting the polynucleotide. The polynucleotides ADM01316-ADM03758 are  
 CC useful in gene therapy, for developing a diagnostic marker or medicines  
 CC for regulating their expression and activity, or as a target of gene  
 CC therapy. The proteins ADM03759-ADM06201 encoded by the polynucleotides  
 CC are useful as pharmaceutical agents. The present sequence represents a  
 CC cDNA sequence of the invention.  
 XX Sequence 2238 BP; 668 A; 440 C; 465 G; 665 T; 0 U; 0 Other;  
 SQ Query Match 5.5%; Score 48; DB 11; Length 2238;  
 Best Local Similarity 50.4%; Pred. No. 0.0018; Mismatches 115; Indels 0; Gaps 0;  
 Matches 117; Conservative 0; Gaps 0;  
 Qy 540 ATATCTTACTTGGAAACAGATGACTGAAATATGACTGCTGAACTGATGCCAGTCCTTCATCA 599  
 Db 637 ATTCTGGATGCCAATGACATTAGCTGATGCCAACTGCTGATGCCAACTGCTGATCA 696

RESULT 9  
 ADA24507  
 ID ADA24507 standard; cDNA; 2887 BP.  
 XX  
 AC ADA24507;  
 XX DT 20-MCN-2003 (first entry)  
 XX DE Human cDNA differential expressed in adipose tissue, INCYTE138185-7.  
 XX KW  
 XX SS; differential expression; adipose tissue; cytostatic; hypotensive;  
 KW ant arteriosclerotic; antidiabetic; anorectic; gene therapy;  
 KW peroxisome proliferator-activated receptor gamma; pPargamma;  
 KW diabetes mellitus; obesity; hypertension; atherosclerosis; prostate cancer;  
 KW prostate cancer; colon cancer; polycystic ovarian syndrome.  
 OS Homo sapiens.  
 XX PN US2003095272-A1.  
 XX PD 22-MAY-2003.  
 XX PP 29-JUL-2002; 2002US-00208408.  
 XX PR 30-JUL-2001; 2001US-0308869P.  
 XX PA (INCY-) INCYTE GENOMICS INC.  
 XX PI Schbaye XM;  
 XX WPI: 2003-606416/57.  
 XX New combination comprising several cDNAs, useful for preparing a  
 PT composition for diagnosing or treating diabetes mellitus, obesity,  
 PT hypertension, atherosclerosis, or cancer of the breast, prostate or  
 PT colon.  
 XX PS Claim 1; Page 47-48; 84pp; English.  
 XX Then invention relates to a new combination comprising 55 cDNAs (ADA24485  
 CC -ADM24539) or their complements that are differentially regulated in an  
 CC adipose sample. Also included are detecting differential expression of  
 CC one or more cDNAs in a sample containing nucleic acids, screening several  
 CC molecules or compounds to identify a ligand that specifically binds a  
 CC cDNA, a vector comprising the cDNA, a host cell containing the vector,  
 CC producing a protein, screening several molecules or compounds, producing  
 CC an antibody and the isolated antibody. The cDNAs comprise sequences which  
 CC are upregulated or downregulated in response to peroxisome proliferator-  
 activated receptor gamma (pPargamma) agonist. The combination comprising  
 CC several cDNAs is useful for preparing a composition for diagnosing or  
 CC cancer of the breast, prostate or colon, or polycystic ovarian syndrome.  
 XX SQ Sequence 2887 BP; 872 A; 552 C; 560 G; 903 T; 0 U; 0 Other;  
 Query Match 5.5%; Score 48; DB 9; Length 2887;  
 Best Local Similarity 50.4%; Pred. No. 0.002; Mismatches 115; Indels 0; Gaps 0;



polynucleotides and polypeptides are useful in gene therapy, vaccines or peptide therapy. The polypeptides have various cytokine-like activities, e.g. stem cell growth factor activity, haemopoiesis regulating activity, tissue growth factor activity, immunomodulatory activity and activin/inhibin activity and may be useful in the diagnosis and/or treatment of cancer, leukaemia, nervous system disorders, arthritis and inflammation. Note: Records for SBQ ID NO 2110 (AARS2581), 2111 (AARS2582) and 3666 (AM80020) are omitted as the relevant pages from the sequence listing were missing at the time of publication
Sequence 4318 BP; 1319 A; 769 C; 814 G; 1416 T; 0 U; 0 Other;
Query Match 5.5%; Score 48; DB 4; Length 4318;
Best Local Similarity 50.4%; Pred. No. 0 0024;
Matches 117; Conservative 0; Mismatches 115; Indels 0; Gaps 0
540 ATATCTACTTGGAAACAGTATGACTGAAATATGACTGTGAAACGATGCCAGCTCTCATCA 599
697 ATTCTGGATGGCATGAAATGCAATTGCTGAACTCTGCTGCCAAACTGCAT 756
Qy 540
600 TATTCGAATTATGGATGACTCTTGGATGATATTCCACATAATTTCACCTCT 659
757 TGTCAAGGGTGGCCAAAAAAATTCGAACTTGTGATATTCCDAAGAATGACTGGCAT 816
Qy 600
660 CTGGCTTATATCTCATCGTACAGCGTACAGCGCATTATTGAGACTTGTCCCGCGA 719
817 CTGGAGATCCTACTAATGATCATAGTAGGGAGGTACCAATACTGTGCACTGTA 876
Qy 660
720 TCGGACATTATTCATCACTATAAGAACAAATGAACTGTGTCACAATCAA 771
877 TAAGGAGGTGAAATAGCCTATGATGTCAGCCAAAAGACTCACCAGTAA 928
Ddb
RESULT 1.2
AAA578892
AA578892 standard; cDNA; 4318 BP.
XX
AA578892;
XX
22-OCT-2001 (first entry)
XX
Human polynucleotide SBQ ID NO 85.
XX
Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer; peripheral nervous system; neuropathy; central nervous system; CNS; Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic; amyotrophic lateral sclerosis; Shy-Drager Syndrome; Chemotactic; chemokinetic; thrombolytic; drug screening; arthritis; inflammation; leukaemia; SS.
XX
Homo sapiens
XX
WO200153312-A1.
XX
26-JUL-2001.
XX
26-DEC-2000; 2000WO-US344263.
XX
23-DEC-1999; 99US-00471275.
XX
21-JAN-2000; 2000US-00488725.
XX
25-APR-2000; 2000US-00552317.
XX
20-JUN-2000; 2000US-00598042.
XX
19-JUL-2000; 2000US-00620312.
XX
03-AUG-2000; 2000US-00653450.
XX
14-SEP-2000; 2000US-00662191.
XX
19-OCT-2000; 2000US-00693036.
XX
29-NOV-2000; 2000US-00727344.
HYSEQ INC.
XX
Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D, Zhang J, Yang Y, Xu C, Xue AJ, Wang Z, Wehrman T, Xu C, Dormanac RT, Zhou P, Goodrich R, Dormanac RT, Zhou P, Goodrich R, Dormanac RT,

XX	WPI; 2001-442253/47.
DR	P-BSDB; AAM38726.
XX	Novel nucleic acids and polypeptides, useful for treating disorders such as central nervous system injuries.
PT	Claim 1, SEQ ID NO 85; 10078pp; English.
XX	The invention relates to human nucleic acids (AAI57798-AAI61369) and the encoded polypeptides (AAM3842-AAM42213) with nootropic, immunosuppressant and cytosstatic activity. The polypeptides or polynucleotide in gene therapy. A composition containing a polypeptide or polynucleotide of the invention may be used to treat diseases of the peripheral nervous system, such as peripheral nervous injuries, peripheral neuropathy and localised neuropathies and central nervous system diseases, such as Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager Syndrome. Other uses include the utilisation of the activities such as: Immune system suppression, Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic and thrombolytic activity, cancer diagnosis and therapy, drug screening, assays for receptor activity, arthritis and inflammation, leukemias and C.N.S disorders. Note: The sequence data for this patent did not form part of the printed specification
XX	Sequence 4318 BP; 1319 A; 769 C; 814 G; 1416 T; 0 U; 0 Other;
SQ	Query Match 5.5%; Score 48; DB 4; Length 4318; Best Local Similarity 50.4%; Pred. No. 0.0024; Matches 117; Conservative 0; Mismatches 115; Indels 0; Gaps 0
Qy	540 ATATCTACTTGGAAACAGTATGACTGAATATGACTGTGACTGTGACCTGTCACATCA 599
Db	697 ATTCTGGATGGCATGAAATGACATTAGCTGATTGCAACCTGCTGCCAAAC1GATCAT 756
Qy	600 TAATCGAAATTATTGGATTTGACTCTTGATTGGATAATCCAGATAATTTCACATCATCT 659
Db	757 TGTCAAGGGTGGCaaaaATATGGCACTTGTGATATTCCAAAAGAATATGACTGGCAT 816
Qy	660 CTCGGCTTATATCCTACTCGCATACGGCAGCATTTATGAGATGTTGTCGGCGGA 719
Db	817 CTGGAGATACCTTAACTAACTGATACAGTACAGTGGAAAGAGTTACCCATACCTGTCAGTGA 876
Qy	720 TCAAGGACATTATTCATCACTATAAGAACAAATCAATCTGTTCAAAATCAA 771
Db	877 TAAGGGTGGAAATAGCATATAGTGTAGTAGGCCAAAAGACTCNCACAAAGTAA 928
RESULT 13	
ACF12844	ACF12844 standard; cDNA; 4318 BP.
XX	
AC	ACF12844;
XX	
DT	10-SEP-2003 (first entry)
XX	
DE	Human cervical cancer cell marker encoding cDNA SEQ ID NO:33.
XX	
KW	Human; cervical cancer; cervical cancer marker; cancer therapy;
XX	
KW	detection; gene therapy; vaccine; gene; ss.
OS	
XX	Homo sapiens.
PN	WO2002101075-A2;
XX	
PD	19-DEC-2002.
XX	
PF	12-JUN-2002; 2002WO-US018638.
XX	
PR	13-JUN-2001; 2001US-0298155P.
PR	13-JUN-2001; 2001US-0298155P.
PR	14-NOV-2001; 2001US-03359365.
XX	

PA (MILL-) MILLENIUM PHARM INC.  
 XX  
 PI Schlegel R, Chen Y, Zhao X, Monahan JE, Kamathkar S;  
 PI Gannavarapu M, Glatt K, Hoersch S;  
 XX  
 WPI; 2003-156967/15.  
 DR P-PSD; ABR92063.  
 XX  
 PT New isolated nucleic acid molecule useful for detecting, characterizing,  
 PT preventing and treating human cervical cancers, in various prognostic and  
 PT diagnostic assays, in pharmacogenomics and in monitoring clinical trials.  
 XX  
 PS Claim 4: Page 160-161; 386pp; English.  
 XX  
 CC ACP12828 to ACP12947 encode the human cervical cancer marker proteins (I)  
 CC given in ABR92067 to ABR92164. A higher level of expression of (I) than  
 CC normal indicates the presence of cervical cancer. Also described: (1) a  
 CC vector (II) containing (I); (2) a host cell (III) containing (I); and (3)  
 CC assessing (M1) whether a patient is afflicted with cervical cancer,  
 CC comprising comparing the level of expression of a marker in patient's  
 CC sample, and the normal level of expression of the marker in a control non-  
 CC cervical cancer sample, where a significant increase in the level of  
 CC expression of the marker in the patient's sample relative to that in the  
 CC control sample is an indication that the patient is afflicted with  
 CC cervical cancer. (I) has cytostatic activity, and can be used in gene  
 CC therapy and in vaccines. (II) is useful in detecting, characterising,  
 CC preventing and treating human cervical cancers. (I) may also be used in  
 CC various prognostic and diagnostic assays, pharmacogenomics and in  
 CC monitoring clinical trials

XX Sequence 4318 BP; 1315 A; 764 C; 832 G; 1407 T; 0 U; 0 Other;  
 XX Sequence 4318 BP; 1315 A; 764 C; 832 G; 1407 T; 0 U; 0 Other;  
 XX Sequence 4318 BP; 1315 A; 764 C; 832 G; 1407 T; 0 U; 0 Other;

Query Match 5.5%; Score 48; DB 8; Length 4318;  
 Best Local Similarity 50.4%; Pred. No. 0.0024;  
 Matches 117; Conservative 0; Mismatches 115; Indels 0; Gaps 0;

Query 540 ATATCTACTTGAAACAGTATGACTGAAATGACTGTGAACTGTGAACCTGCACTCTCATCA 599  
 Db 728 ATTTCGGATGGCAATGAAATGACTGTGAACTGTGAAATGACTGTGAACCTGCAACTCTCATCA 599  
 Query 600 TATTGGAATTATTGGATTGTCACTTCTGGATTGATATTGAGATTCACATAATTCTACTCATCT 659  
 Db 788 TGTCAGGTGGTGGCAAAATAATGCAATTGCACTTGTGAACTTGTGAAATGACTGTGAACTGTGCA 659  
 Query 660 CTGGGCTTATATCCTCATGCTACCCATACGAGATTTGAGATTTGAGATTTGCCCCCGA 719  
 Db 784 CTGGAGATACCTAACATGCTACAGTAGGAGCTACAGTAGGAGCTACCATATCCTGCAATCTGCA 719  
 Query 720 TCAGGACATTATTCACTCATATAAGAACAAATGATCTGTCAAAATCA 771  
 Db 908 TAAGGGGTTGAAATAGCATATAGGTGAGCTACAGTAGGAGCTACCATATCCTGCAATCTGCA 959

RESULT 15  
 ADP21232 ID ADP21232 standard; cDNA; 4318 BP.  
 AC AC ADP21232;  
 XX AC ADP21232;  
 XX DT 18-NOV-2004 (first entry)  
 DE PRO polypeptide encoding cDNA SEQ ID NO:326.  
 XX KW ss; Gene; PRO; antiinflammatory; antiarthritic; antirheumatic;  
 KW immunosuppressive; osteopathic; antiidiabetic; dermatological;  
 KW antipsoriatic; antiallergic; antiasthmatic; hepatotropic; respiratory;  
 KW gene therapy; immune system.  
 XX OS Unidentified.  
 XX OS  
 XX PN WO2004041170-A2.  
 XX PD 21-MAY-2004.  
 XX PN WO2004065545-A2.  
 XX PD 05-AUG-2004.  
 XX PN 15-JAN-2004; 2004WO-US00110.0.  
 XX PR 01-NOV-2002; 2002US-0423394P.

RESULT 14  
 ADR25580 ID ADR25580 standard; DNA; 4318 BP.  
 AC AC ADR25580;  
 XX DT 21-OCT-2004 (first entry)  
 DE Breast cancer prognosis marker #1441.  
 XX KW ds; breast cancer; prognosis; gene expression; diagnosis.  
 XX OS Homo sapiens.  
 XX PN WO20040410655-A2.  
 XX PD 30-OCT-2003; 2003WO-US034312.  
 XX PR 01-NOV-2002; 2002US-0423394P.

XX (GETH ) GENENTECH INC.  
 PA  
 XX Clark H, Schoenfeld J, Van Lookeren M, Williams PM, Wood WI;  
 PI  
 PI Wu TD;  
 XX  
 DR WPI: 2004-419628/39.  
 DR P-PSDB; ADB23233.

PT New PRO polypeptides and polynucleotides, useful for treating e.g.  
 PT erythematosus, rheumatoid arthritis, diabetes mellitus, immune-mediated  
 PT renal disease, or demyelinating diseases of the central or peripheral  
 PT nervous system.

XX

PS Claim 1; SEQ ID NO 326; 2940pp; English.

XX The invention relates to a novel isolated nucleic acid and the PRO  
 CC polypeptide encoded by it. A protein of the invention has  
 CC antiinflammatory, antiarthritic, antirheumatic, immunosuppressive,  
 CC osteopathic, antidiabetic, dermatological, antipruritic, antiallergic,  
 CC antiinflammatory, hepatotropic, and respiratory activity. A polynucleotide,  
 CC polypeptide, or antibody that specifically binds to the  
 CC agonist, antagonist, or antibody that specifically binds to the  
 CC polypeptide is useful for treating an immune related disorder such as  
 CC systemic lupus erythematosus, rheumatoid arthritis, osteoarthritis,  
 CC juvenile chronic arthritis, a spondyloarthropathy, systemic sclerosis, an  
 CC idiopathic inflammatory myopathy, Sjogren's syndrome, systemic  
 CC vasculitis, sarcoidosis, autoimmune haemolytic anaemia, autoimmune  
 CC thrombocytopenia, thyroiditis, diabetes mellitus, immune-mediated renal  
 CC disease, a demyelinating disease of the central or peripheral nervous  
 CC system, idiopathic demyelinating polyneuropathy, Guillain-Barre syndrome,  
 CC a chronic inflammatory demyelinating polyneuropathy, a hepatobiliary  
 CC disease, infectious or autoimmune chronic active hepatitis, primary  
 CC biliary cirrhosis, granulomatous hepatitis, sclerosing cholangitis,  
 CC inflammatory bowel disease, gluten-sensitive enteropathy, Whipple's  
 CC disease, an autoimmune or immune-mediated skin disease, a bullous skin  
 CC disease, erythema multiforme, contact dermatitis, psoriasis, an allergic  
 CC disease, asthma, allergic rhinitis, atopic dermatitis, food  
 CC hypersensitivity, urticaria, an immunologic disease of the lung,  
 CC eosinophilic pneumonia, idiopathic pulmonary fibrosis, hypersensitivity  
 CC pneumonitis, a transplantation associated disease, graft rejection or  
 CC graft-versus-host disease. The present sequence encodes a PRO protein of  
 CC the invention.

XX Sequence 4318 BP; 1315 A; 764 C; 832 G; 1407 T; 0 U; 0 Other;

Query	Match	Score	Length
Qy	540 ATATCTACTTGGAAACAGTATGACTGAATATGACTGTGAACCTGATGCACCTCTTCATCA	599	4318
Db	728 ATTCTGGATGGATGATGATGCACTTGGATTCATGACCTGATTCGAACTGTGCTGCCAACTGATAT	787	
Qy	600 TATTCGAATTATTGGATTTGCTTGTCACTTCTGGATTCGATACTTCAATTTCACTCTATCT	659	
Db	788 TGTCAAGGGGGTCCAAAAAAATGGAACTTGTGATTCGAAATGGATTCAGAAATGACTGGCAT	847	
Qy	660 CTGGCTTATATCTCATGCTACCGTACAGAGCTTATTGAGAGTTGTCGGCGGA	719	
Db	848 CTGGAGATACCTAACATGCTACAGTGGGAGCTACCAATCTGTCCTGAGTGA	907	
Qy	720 TCAGGACATTATTGATCACTATAAAAGACAAATGAAATGTTGACAAATGAGTCAACATCAA	771	
Db	908 TAAGGAGGTGAAATAGGATATAGTGTAGCCAAAAGACTCACCAGTAA	959	